NAVAL SUPPORT ACTIVITY MID-SOUTH

A-A SEQUENTIAL REMEDIATION TREATABILITY STUDY REPORT AOC A — NORTHSIDE FLUVIAL GROUNDWATER

CTO-0094

Prepared for:



Department of the Navy
Southern Division
Naval Facilities Engineering Command
North Charleston, South Carolina

Prepared by:



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May 17, 2002

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ABBREVIATIONS AND ACRONYMS

a-a anaerobic-aerobic AOC Area of Concern

BCT BRAC Cleanup Team bls below land surface

BRAC Base Closure and Realignment Act

BTEX Benzene, Toluene, Ethylbenzene, Xylene

CFUs colony forming units

CMS Corrective Measures Study

CO₂ carbon dioxide

CRP Community Relations Plan

CSI Confirmatory Sampling Investigation

DCA dichloroethane
DCE dichloroethene
DO dissolved oxygen
DPT direct push technology

E/A&H EnSafe/Allen & Hoshall EIC Engineer-In-Charge

gpm gallons per minute

HPC heterotrophic plate count

HSWA Hazardous and Solid Waste Amendments

ID inside diameter

IDW investigation-derived waste IRP Installation Restoration Program

K_{xy} hydraulic conductivity

kg kilogram

lbs pounds

MCLs maximum contaminant levels

mg/L milligram per liter

MNA monitored natural attenuation

mV millivolts

NSA Naval Support Activity

ABBREVIATIONS AND ACRONYMS (continued)

O&M operations and maintenance ORP oxidation-reduction potential

PCE tetrachloroethene

psi pounds per square inch

PVC polyvinyl chloride

QAPP Quality Assurance Project Plan

RAB Restoration Advisory Board RBCs risk-based concentrations

RCRA Resource Conservation and Recovery Act

RFA/RFI RCRA Facility Assessment/RCRA Facility Investigation

SAP Sampling and Analysis Program SWMU solid waste management unit

TCE trichloroethene

TDEC Tennessee Department of Environment and Conservation

TKN total kjeldahl nitrogen
TOC total organic carbon
TS treatability study

 μ g/kg micrograms per kilogram μ g/L micrograms per liter

USEPA United States Environmental Protection Agency

USGS United States Geological Survey

VC vinyl chloride

VOCs volatile organic compounds

Section 1: Introduction

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1.0 INTRODUCTION

An anaerobic-aerobic (A-A) sequential remediation treatability study of volatile organic carbon

(VOC) contamination in the fluvial deposits aquifer located on part of the former Northside of

Naval Support (NSA) Mid-South, Millington, Tennessee was conducted by EnSafe Inc. from

March to December 2000. This A-A treatability study is based on the Corrective Measures Study

(CMS) Work Plan (EnSafe, 2000) for Area of Concern (AOC) A (the Northside fluvial

groundwater), and methods presented in the AOC A — Northside Fluvial Groundwater A-A

Sequential Remediation Treatability Study Work Plan (EnSafe, 1999).

The A-A sequential treatment technology is based on enhancing the biodegradation of VOCs that

are present in the aquifer beneath AOC A, as described in Section 2 of this report. The

primary purpose of the study was to determine the feasibility of using A-A sequencing to

remediate the AOC A groundwater plume; specifically, the area of higher chlorinated solvent

concentrations in the vicinity of monitoring well 007G04LF. A secondary purpose was to

determine the chemical, biochemical, and physical impact of the treatability study on the aquifer.

Five wells were installed for the treatability study in February 2000 near existing wells 04LF and

04UF. Two additional monitoring wells were installed in August 2000. The wells installed for

the treatability study include:

• One 4-inch inside diameter (ID) polyvinyl chloride (PVC) extraction well (57LF)

• Two 4-inch ID PVC reinjection wells (60LF, 61LF)

• Four 2-inch ID PVC monitoring wells (58LF, 59LF, 62LF, 63LF)

Each well was installed in the fluvial deposits, the top of which occurs at approximately 30 feet

below land surface (bls) and extends to 75 feet bls in the treatability-study area. The extraction

and reinjection wells each have 30-foot-long PVC screens for the system at the bottoms. The

pumps, piping, instrumentation, tanks, electrical wiring, and housing were installed in February 2000, as described in the work plan (EnSafe, 1999). Section 5 of this report provides more information on well installation and the system setup.

The system operated from March 14, 2000 to December 15, 2000. During this time, periodic field monitoring data and monthly analytical data were collected, as described in Section 6 of this report. Results of the study are presented and analyzed in Section 7 of this document.

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2.0 REMEDIAL TECHNOLOGY DESCRIPTION

2.1 Introduction

A-A sequential groundwater treatment, also known as "two-zone interception treatment," is

designed for enhanced in-situ bioremediation of chlorinated solvent contamination in groundwater.

The USEPA has demonstrated the treatment as an emerging technology under the

Superfund Innovative Technology Evaluation (SITE) program.

2.2 Theory

Most chlorinated solvents at contaminated groundwater sites are amenable to biodegradation.

However, compared with petroleum hydrocarbons, chlorinated solvents are more sensitive to

groundwater oxidation-reduction potential (ORP), availability of natural organic carbon or

anthropogenic organic substrates (e.g., benzene, toluene, ethyl benzene, and xylene [BTEX] or

other man-made carbon sources), and natural groundwater electron acceptors such as

dissolved oxygen, nitrate, sulfate, and carbon dioxide.

While petroleum hydrocarbons can serve as a primary organic substrate (food source that provides

energy) or electron donor for microorganisms, chlorinated solvents — particularly the

highly chlorinated solvents such as perchlorethylene (PCE) and trichloroethylene (TCE) — are not

a direct food or energy source. PCE and TCE serve more as electron acceptors much as oxygen,

nitrate, sulfate, and carbon dioxide do in BTEX or natural organic carbon degradation. In

other words, anaerobic (absence of dissolved oxygen) or reduced conditions in an aquifer are more

suitable to PCE and TCE degradation. Moreover, the more strongly reduced an aquifer is, the

more readily PCE and TCE degrade.

The lesser chlorinated solvents such as 1,2-dichloroethylene (1,2-DCE) and vinyl chloride (VC) are more likely to serve as organic substrates (electron donors) or co-substrates and are more amenable to biodegradation in the presence of oxygen. These solvents are breakdown products (daughter compounds) of PCE and TCE degradation.

2.3 Treatment Process

The degree of anaerobicity or aerobicity of and aquifer can be estimated from redox measurements of the groundwater it contains. The lower the redox potential (measured in millivolts) of an aquifer, the more anaerobic or strongly reducing it is. In general, redox potentials less than +50 millivolts (mV) represent anaerobic (reducing) conditions. If redox measurements near a PCE and/or TCE plume are greater than +50 mV, nutrients (nitrate and phosphate compounds) and substrate (organic carbon) can be added to consume oxygen and drive the system to more strongly anaerobic or reducing conditions. Generally, enough carbon is added to create anaerobic conditions and be available as a food source while highly chlorinated solvents such as PCE and TCE are subsequently degraded. Conversely, if redox measurements near a 1,2-DCE and/or VC plume are less than +50 mV, air sparging of the aquifer can be used to increase oxygen availability and allow maximum biological consumption of substrates such as 1,2-DCE and VC.

2.4 Methodology

Creating an anaerobic zone upgradient of an aerobic zone within a PCE/TCE-contaminated groundwater plume establishes a sequential A-A zone that can degrade these compounds sequentially to innocuous gaseous end products. Moreover, flow through these zones can be accelerated by installing low-flow extraction wells downgradient of the aerobic zone and reinjecting the extracted groundwater, which has ben amended with carbon and other nutrients as described below, upgradient of the anaerobic zone.

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Section 2: Remedial Technology Description

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Anaerobic Zone

An anaerobic zone in the contaminated part of the aquifer is created by pumping groundwater from

downgradient extraction wells and adding carbon and other nutrients to it aboveground before

reinjecting the groundwater into upgradient wells. The groundwater is first pumped to an

aboveground chemical amendment system where carbon (fructose or acetate) and nutrients

(ammonium phosphate) are added before the water is reinjected into the aquifer. The carbon and

nutrients provide a ready food source that stimulates microbial respiration, which consumes all the

available oxygen in the treated groundwater.

This recirculation process (extraction and reinjection) continues until an anaerobic zone is

gradually created near the reinjection wells. Highly chlorinated solvents such as PCE and TCE

are amenable to reductive dechlorination (biological removal of the chloride atoms) under

anaerobic conditions. In other words, after the anaerobic zone is established, microorganisms will

turn to sources other than oxygen, such as the higher chlorinated VOCs, for respiration.

Aerobic Zone

Anaerobic reductive dechlorination results in the formation of lesser-chlorinated

daughter products, namely VC and 1,2-DCE. However, these compounds break down more

readily in an aerobic environment. Therefore, if needed, an aerobic zone may be created near the

downgradient extraction wells by injecting air into the aquifer via sparging wells connected to an

aboveground blower. Sparging is generally performed intermittently, based on groundwater

dissolved oxygen (DO) concentrations in area monitoring wells. Carbon and nutrients can also

be added to the air sparging wells to enhance the aerobic degradation of 1,2-DCE and VC.

Aerobic degradation of VC forms innocuous end products such as ethane, ethene, carbon dioxide,

and water.

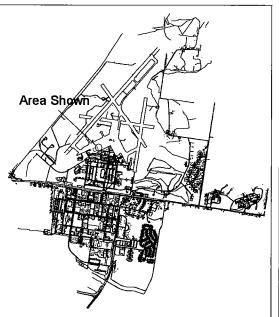
3.0 SITE DESCRIPTION

3.1 Introduction

Twelve solid waste management units (SWMUs) were identified on the NSA Mid-South Northside for Resource Conservation and Recovery Act (RCRA), Confirmatory Sampling Investigation (CSI) or RCRA Facility Investigation (RFI) characterization. During the November 1994 direct push technology (DPT) groundwater screening investigation for the SWMU 7 (N-126 plating shop drywell; now part of AOC A) RFI, chlorinated solvents (e.g., TCE and DCE) were detected in groundwater in the fluvial deposits aquifer. As the area of investigation expanded while the nature and extent of contamination were being defined, it became apparent that groundwater contamination in the airfield apron area was widespread and that SWMU 7 was not the primary source. The focus of the SWMU 7 groundwater investigation then shifted from the dry well to the entire airfield apron area, and ultimately to the entire NSA Mid-South Northside, as scattered pockets of contaminated groundwater in the fluvial deposits were identified. As a result, the Base Closure and Realignment Act (BRAC) Cleanup Team (BCT) decided to take a "holistic" approach to the Northside groundwater investigation and any subsequent CMS, creating AOC A, the Northside fluvial groundwater, to be evaluated as one unit rather than on a site-by-site basis.

Although the fluvial deposits groundwater beneath a large part of the Northside is included in AOC A, the CMS will focus on three main areas: (1) the plume areas where the highest VOC contaminant concentrations have been detected on the east side of Building N-126 near monitoring well 007G04LF, (2) the area once occupied by former Building N-6, and (3) the plume area north of the main runway which appears to extend offsite. The treatability study described in this report was conducted in the area east of Building N-126 where the highest chlorinated solvent concentrations have been identified in the groundwater in the fluvial deposits. A site map of the treatability study area is included as Figure 3-1.







- **Extraction Well**
- Injection Well Monitoring Well
- Road Building



Figure 3-1 Site Map of Study Area NSA Mid-South

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3.2 Site Geology and Hydrogeology

The fluvial deposits beneath the former NSA Mid-South airfield apron area and throughout most

of the Memphis area are made up of poorly sorted sand and gravel of Pleistocene to possibly

Pliocene age, with minor amounts of clay as interstitial material, and occasional clay lenses

generally no more than a few inches thick. Fine to medium sand in the upper sections coarsens

with depth. Gravel occurs as lenses at various horizons in the fluvial deposits, but is more

common in the lower part of the unit. The thickness of the fluvial deposits, which are

fully saturated in the treatability study area, ranges from 26 to 64 feet within AOC A.

The fluvial deposits are overlain, and water in the formation is confined or semiconfined in by

Pleistocene-age loess, a relatively low-permeability unit of silt and clayey silt that ranges from

25 to 45 feet thickness within AOC A. A perched groundwater zone occurs in the loess

throughout much of NSA Mid-South and varies from 4 to 8 feet bls. However, this

perched groundwater zone is absent beneath much of the apron where recharge is inhibited by the

large area of concrete pavement.

The base of the fluvial deposits (ranges from about 70 to 100 feet bls) lies uncomformably on top

of the Cockfield Formation of Eocene age, which, together with the underlying Cook Mountain

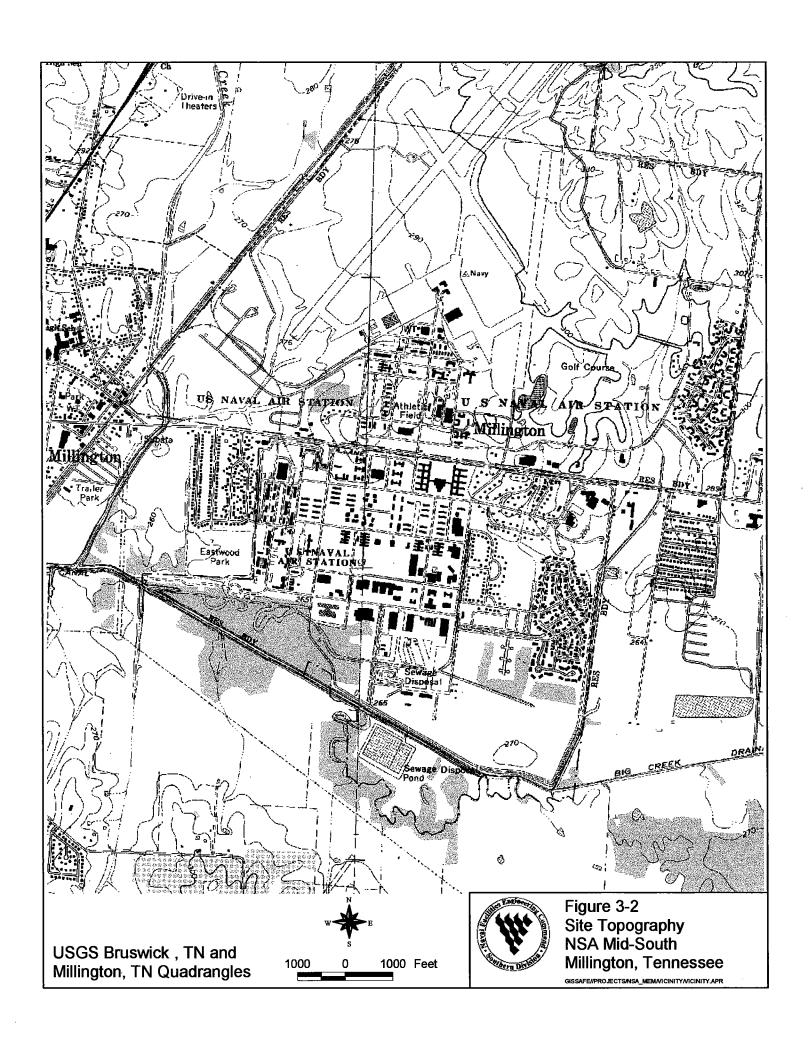
formations (upper units of the Claiborne Group), forms the lower confining unit for the

fluvial deposits and upper confining unit for th Memphis aquifer in the Memphis area.

Water levels in monitoring wells screened in the Cockfield Formation, which is composed

primarily of fine sand and silt with interbedded clay in the NSA Mid-0South area, are also

confined by clay beds in the formation and are nearly equal to those in the fluvial deposits.



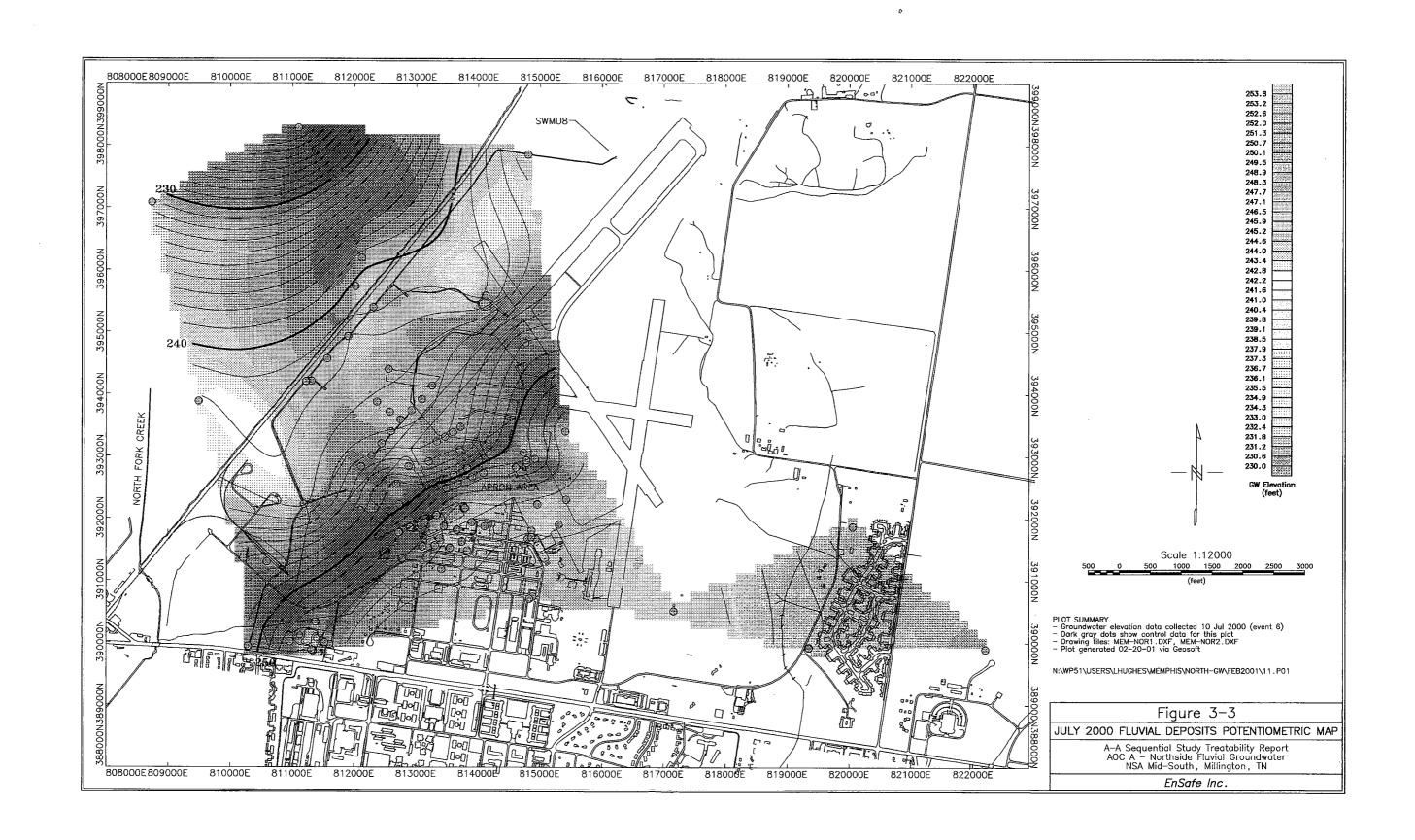
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Section 3: Site Description

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The Cook Mountain Formation, which contains the most aerially extensive clay in the upper part of the Claiborne Group in Memphis and Shelby County, serves as primary components of the confining unit separating groundwater in the fluvial deposits and Cockfield Formation from groundwater in the Memphis aquifer. The Cook Mountain Formation at NSA Mid-South consists predominantly of clay and silt; however, minor lenses of silty fine sand may be present locally. Geophysical logs from public supply wells indicate the Cook Mountain Formation ranges from 10 to 60 feet thick at NSA Mid-South (Carmichael et al., 1997). A topographic map of the NSA Mid-South Northside is shown on Figure 3-2.

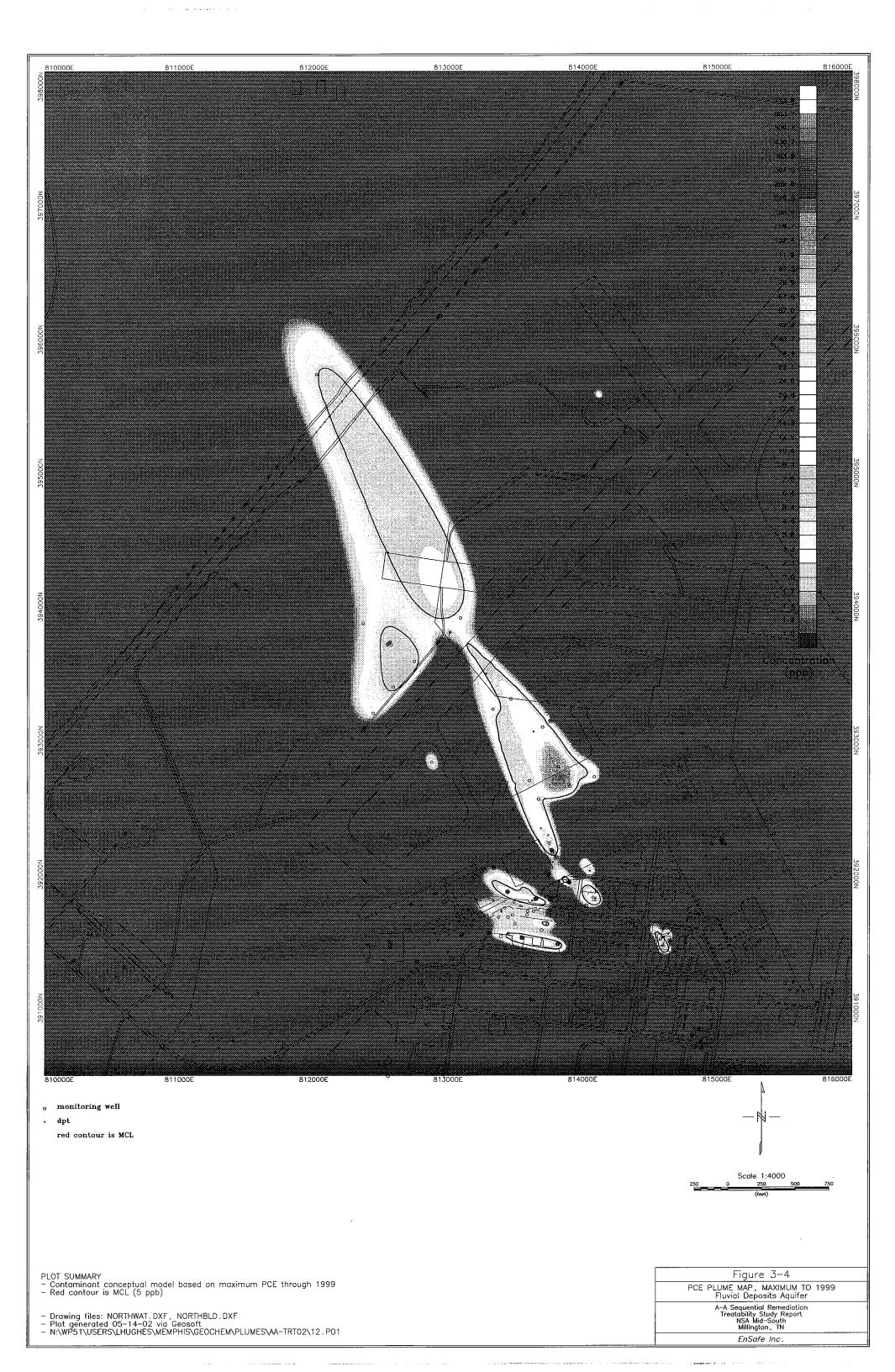
In July 2000, EnSafe measured groundwater elevations in existing Northside monitoring wells screened in the fluvial deposits. Figure 3-3 shows groundwater in this unit flowing radially away from a south-to-northwest trending ridge. Potentiometric data from the apron area indicate that groundwater in the fluvial deposits is semiconfined to confined and flows north and west with an average hydraulic gradient of 0.004 to 0.008 foot per foot. Results of an aquifer test in the fluvial deposits southwest of the apron area produced an estimated horizontal hydraulic conductivity (K_{xv}) of 5.3 feet per day (Robinson et al., 1997), which yields a groundwater velocity from 31 to 62 feet per year, using a 25% assumed effective porosity value and the hydraulic gradients listed above. Likewise, another aquifer test in the fluvial deposits conducted north of the main runway produced an estimated K_{xy} of 59 feet per day. Using this K_{xy} value with the flatter gradients north of the runway (0.0017 feet per foot) and the same effective porosity, the groundwater velocity in this area is approximately 145 feet per year.

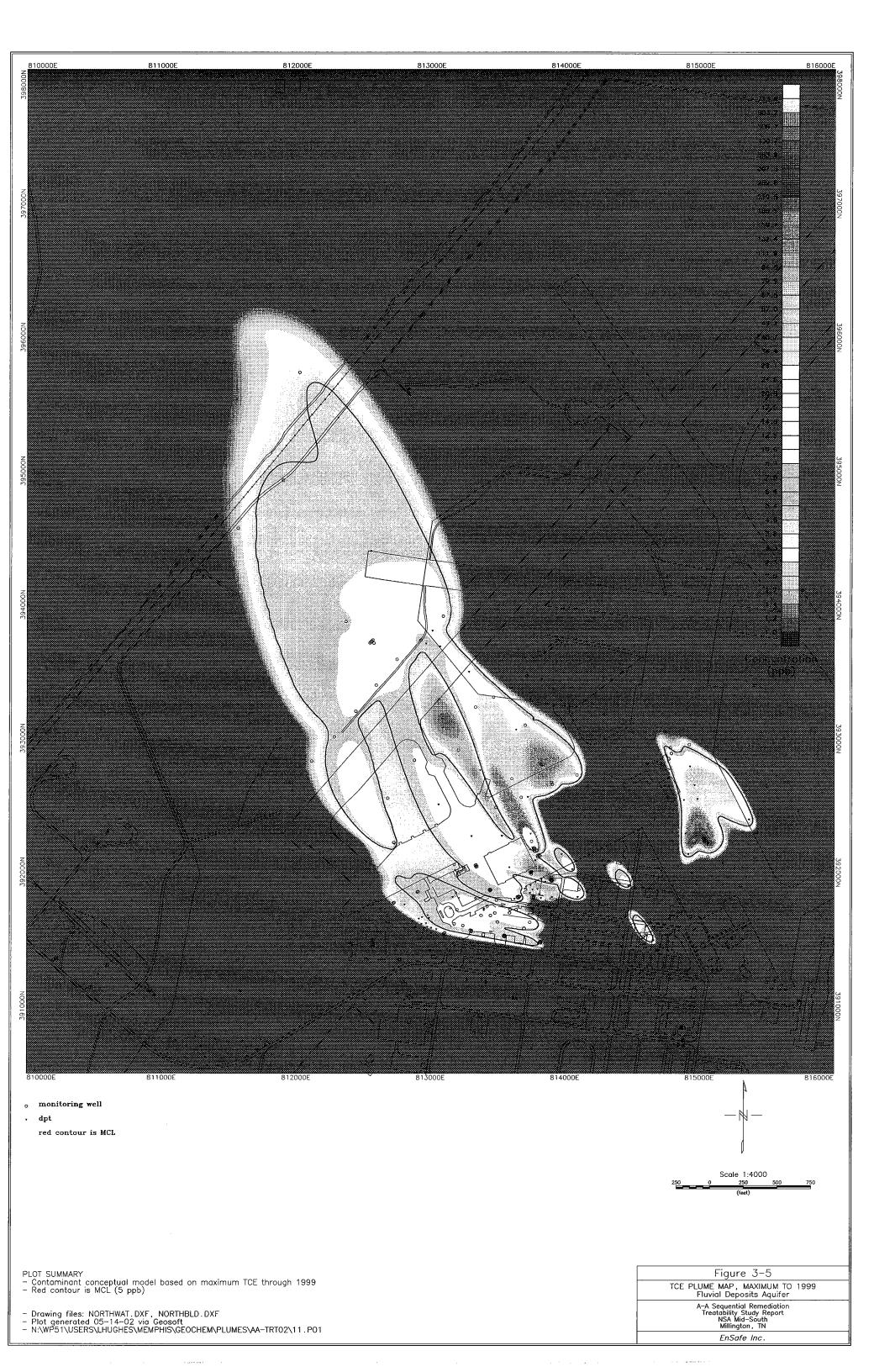


3.3 Nature and Extent of Contamination

A primary reason for designating the Northside fluvial deposits groundwater as an AOC was to expedite the CMS process through collectively evaluating all the SWMUs or contaminant source areas to the fluvial deposits groundwater in this area. The apron-area RFI showed that numerous areas containing multiple VOCs are present beneath the apron at concentrations exceeding maximum contaminant levels (MCLs), and thus warranting corrective measures. The RFI report and addendum for AOC A (EnSafe, 1998 and 2000) present all the fluvial deposits data collected in the apron area through July 1999. Data collected after July 1999 are presented in various technical EnSafe memorandum prepared for the BCT.

The fluvial-deposits data set is large and cumbersome because of multiple SWMUs, multiple sampling events with varying analytical suites, and the number of monitoring wells completed in three zones within the fluvial deposits (upper, middle, and lower parts). Primary contaminants of concern identified in the fluvial deposits are PCE, TCE, 1,2-DCE, 1,1-DCE, 1,2-dichloroethane (DCA) 1,1-DCA, carbon tetrachloride, chloroform, VC, and benzene. Because analytical summary tables for this data set are lengthy and are presented in the AOC A RFI report (EnSafe, 1998), they will not be included in this document. Since most of the significant contamination is from PCE, TCE, and their various daughter products, the treatability study monitored these solvents primarily. Figures 3-4 and 3-5 show the interpreted plumes for PCE and TCE, respectively, according to data collected during the RFI.





Section 4: Treatability Study Objectives

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4.0 TREATABILITY STUDY OBJECTIVES

4.1 Primary Objectives

The main objective of this study was to determine the feasibility of using the A-A sequential technology to degrade and remediate the chlorinated VOC groundwater plumes in the fluvial deposits at AOC A. Although this technology is based on strong microbial principles, it has seen successful application only recently and is still considered an innovative technology. Furthermore, its feasibility often depends on site-specific chemical, geological, and hydrogeological variabilities that are difficult to reproduce in a laboratory. Therefore, a pilot-scale treatability study was needed to assess its effectiveness at AOC A.

The treatability study focused on the plume in the area of monitoring well 007G04LF. Study results will be used to compare this technology with other treatment alternatives and to provide cost and design data for full-scale implementation if this technology is selected as a final remedy or part of the final remedy for this site.

4.2 Secondary Objectives

Secondary objectives of the treatability study included evaluating field parameters necessary to monitor system operation such as DO, pH, oxidation-reduction potential (ORP), heterotrophic plate counts, nutrients, and total organic carbon (TOC). Changes in the values of these parameters (particularly DO) and the time period in which these changes occur can be used to determine how effectively groundwater is being amended to degrade chlorinated solvents. Another secondary objective was to monitor groundwater extraction and reinjection rates and changes in groundwater levels in the area monitoring wells to help evaluate groundwater recirculation patterns in the test area.

5.0 TREATABILITY SYSTEM SET-UP

5.1 System Elements

The A-A sequential system included the extraction well, reinjection wells, monitoring wells, and

an aboveground groundwater chemical amendment system. The equipment layout is shown on

Figure 5-1, while the system process and instrumentation diagram is shown on Figure 5-2.

Groundwater Extraction Well

The 4-inch diameter PVC groundwater extraction well 007G57LF, which is screened in the

fluvial deposits, was installed in December 1999. It is approximately 75 feet deep with a

30-foot-long, 0.02-inch slotted PVC screen from 42 to 72 feet bls. The well had a

pneumatic submersible pump capable of pumping up to 7 gallons per minute (gpm), which was

connected by a 3/4-inch hose to an aboveground chemical feed tank. Groundwater was pumped

from this well to a 500-gallon aboveground holding tank or a 100-gallon chemical mixing tank.

Groundwater Reinjection Wells

Groundwater from the extraction well was amended with nutrients and substrate in the

aboveground system. Amended water was then reinjected by gravity flow into the aquifer via

two 4-inch diameter PVC wells (007G60LF and 007G61LF) screened in the fluvial deposits which

were installed in November/December 1999. They were approximately 75 feet deep and were

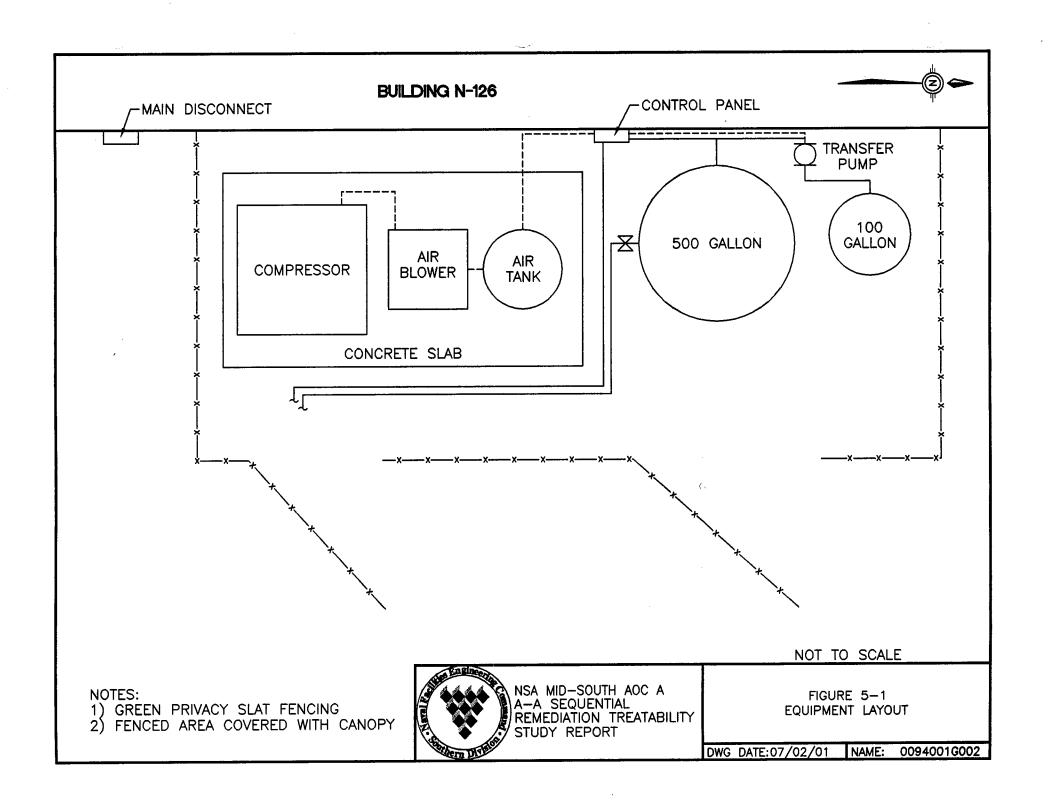
screened with 0.02-inch slotted PVC screen, 30 feet long, from 45 to 75 feet bls.

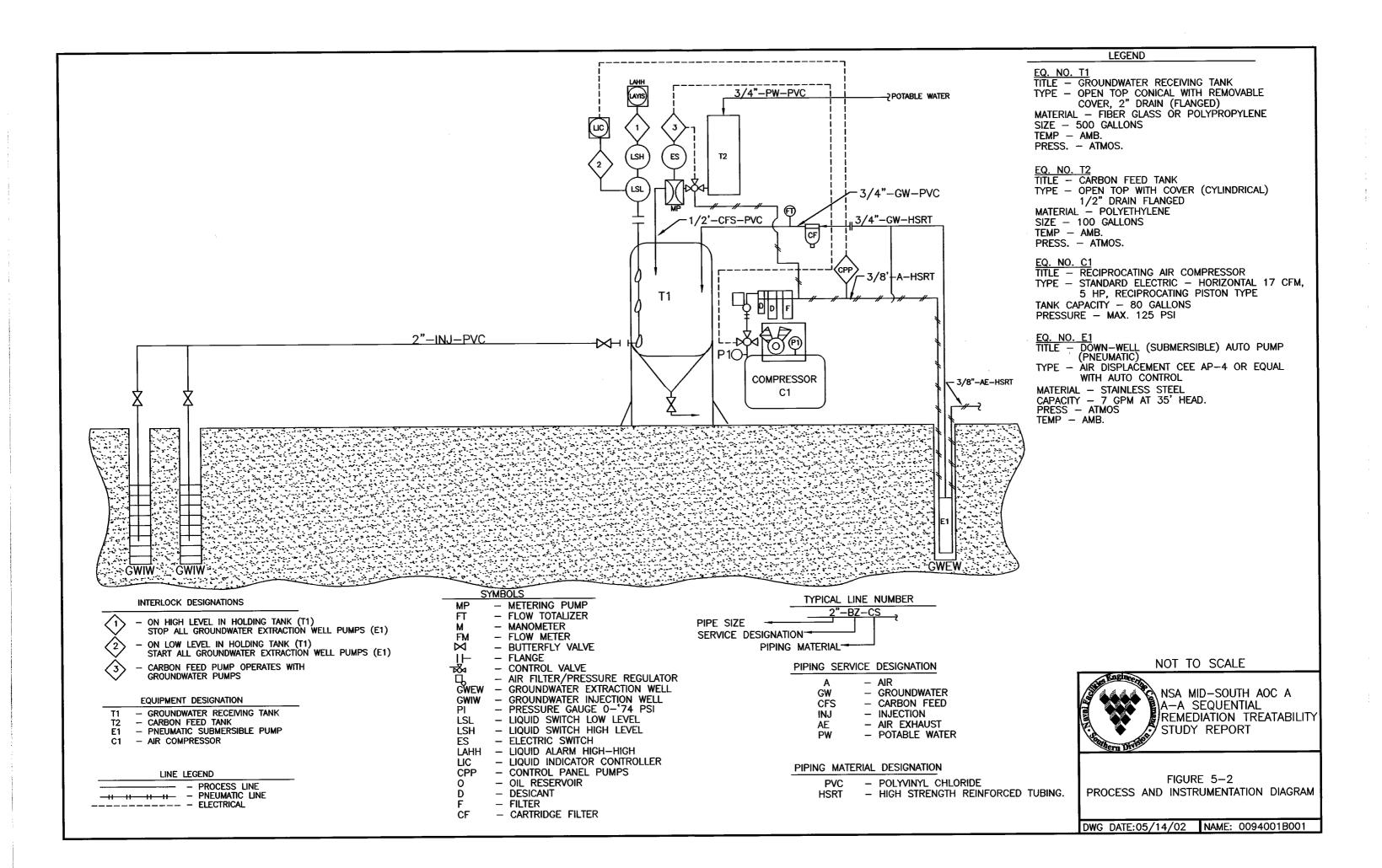
Treatability Study Monitoring Wells

In addition to the four existing monitoring wells (007G03LF, 007G04LF, 007G04UF, and

007G21LF), two 2-inch-diameter PVC wells (007G58LF and 007G59LF) were installed

November 1999 to monitor the progress of the treatability system.





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Section 5: Treatability System Set-Up

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They were approximately 75 feet deep and screened in the fluvial deposits from 42 to 72 feet bgs

with 0.02-inch slotted PVC screen. In August 2000, two additional 2-inch-diameter PVC wells

(007G62LF and 007G63LF) were installed with the same specifications as the other

monitoring wells installed in November 1999 (see Appendix A).

All new monitoring wells were sampled from two depth intervals. The last character of the

sample identification denotes the interval sampled, "A" for the upper part of the fluvial deposits

(47 feet from the top of casing) or "B" for the lower part of the fluvial deposits (5 feet from the

bottom of the well).

Well Specifications

All wells were installed in a boring drilled to a depth that targeted the base of the fluvial deposits.

The wells are flush mounted at ground surface and have a well vault installed over the tops of the

casings. The completed drill holes had outer diameters of 8 to 12 inches, and were large enough

to accommodate a 2- or 4-inch ID well screen and standpipe.

After installation, the wells were thoroughly developed by the drill crew for at least

2 hours each, using a combination of pumping, surging, and flushing with potable water. All

investigation-derived waste (IDW) from drilling and developing the wells were managed as

hazardous waste. Development was completed when the onsite engineer or geologist judged the

well to produce clear water and to be hydraulically responsive.

Submersible Pump

The air-operated submersible pump placed in the extraction well was a Clean Environmental

Equipment (CEE) Standard AP-4 pump that could extract 7 gpm against a total head of 35 feet.

The air supply was regulated by a controller-less total auto pump system. Air requirements,

supplied by the compressor system, were 90 pounds per square inch (psi). To prevent overflow

Section 5: Treatability System Set-Up

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of the 500-gallon holding tank, the autopump system was connected to a tank-full shut-off system

that would block air supply to the submersible pumps if the holding tank became full. Air supply

could also be turned on or off manually by check valves.

Chemical Feed Pump

An air-operated diaphragm pump was used to pump water from the

100-gallon chemical mixing tank to the 500-gallon holding tank. The air supply was regulated by

the same pump system as that used for the submersible pump. The air supply could be turned on

or off manually by check valves.

Compressor System

An electrically powered, rotary screw-type compressor supplied 125 psi that powered

the submersible pumps and chemical-feed diaphragm pump. The three-phase electrical motor

required 230/460 volts at 60 hertz. The compressor was equipped with an air-drying system to

reduce buildup of condensation in the pressure tank.

Groundwater Filter System

A filter system was installed in mid-April to keep fine solids from entering the reinjection wells

and thus clogging their screens. The filter system included a 10/20 micron, cartridge filter and

a pressure gauge to indicate clogging. The filters were replaced weekly or when the pressure

exceeded 15 psi.

Equipment Housing

The blower system, controllerless auto pump system, chemical-feed pump, compressor,

air drying system, 500-gallon holding tank, and 100-gallon chemical mixing tank were housed in

a fenced area with a canopy. The housing provided protection for the equipment and

controlled access to the system.

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Groundwater Recirculation System

The recirculation system was made up of the extraction well, submersible pump, filter system, 500-gallon holding tank, chemical-feed pump, 100-gallon chemical mixing tank, and reinjection wells. The 500-gallon polyethylene holding tank had an inlet at the top and a 4-inch outlet at the bottom. The inlet was connected to the groundwater extraction well via the 3/4-inch hose and a PVC pipe diverted the groundwater influent to the bottom of the tank to avoid aeration and possible volatilization of the VOCs. The outlet was connected to the reinjection wells by an underground PVC piping network system. Water from the holding tank flowed by gravity to the reinjection wells. The 100-gallon chemical mixing tank was adjacent to the 500-gallon holding tank. A 1-inch inlet at the top of the 100-gallon tank allowed groundwater from extraction well 007G57LF to be pumped into the tank where carbon and nutrients were mixed into the groundwater. A chemical-feed diaphragm pump was used to transfer the amended groundwater from the mixing tank into the holding tank.

5.2 Study Area

RFI results for the treatability study area at AOC A show that most of the chlorinated VOC contamination is concentrated in the lower part of the fluvial deposits. TCE is the most prevalent chlorinated solvent detected, followed by carbon tetrachloride and PCE. The highest concentration of any single chlorinated solvent was in a groundwater sample from well 007G04LF, northeast of Building N-126, where TCE was detected at 4,400 micrograms per liter (μ g/L) during the March 25, 1999 RFI sampling event. Because of the high concentrations in the area of monitoring well 007G04LF, the pilot study focused on the plume in this area.

6.0 SUMMARY OF FIELD ACTIVITIES

6.1 System Augmentations

During the pilot study, groundwater was pumped from the extraction well (007G57LF) to the 500-gallon holding tank in the equipment area. Two to three times a week, groundwater from 007G57LF was diverted to the 100-gallon chemical feed tank. When this tank was full, nutrients were added and mixed with the groundwater in the 100-gallon tank. Table 6-1 shows the augmentation schedule.

Table 6-1 Augmentation Schedule (per 100 gallons of water)									
	Period Carbon Source Nutrients								
Start	End	Days	Туре	Quantity (lb)	Type	Quantity (lb)			
3/14/00	3/22/00	8.	fructose	1.1	ammonium phosphate	0.11			
3/22/00	4/5/00	14	fructose	2.2	ammonium phosphate	0.22			
4/5/00	4/26/00	21	fructose	2.2	ammonium phosphate	0.22			
4/26/00	6/26/00	61	fructose	1.1	ammonium phosphate	0.22			
7/7/00	12/8/00	154	sodium acetate	25	ammonium phosphate	0.22			

Note:

lb = pound

During the nine months of the study, 19.3 pounds (lbs) of fructose, 8.7 lbs of ammonium phosphate, and 620 lbs of sodium acetate were added to the system. Typically, 100-gallons of carbon and/or nutrient-enriched groundwater were pumped from the 100-gallon tank into the 500-gallon holding tank. The diaphragm pump was used to pump water from the 100-gallon tank to the 500-gallon tank. More than 1.5 million gallons of water were recirculated during the nine-month study at flow rates ranging from 2 to 10 gpm. As discussed in Section 7.3, well-screen clogging affected re-injection rates.

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6.2 Field Monitoring

Groundwater wells in the test area were monitored for key field parameters such as pH, DO, ORP, and carbon dioxide (CO₂) to optimize system operation and assess the geochemical response of the treatability study during the evaluation process. Baseline field-monitoring data were collected a week before the system was turned on. Wells monitored for baseline field parameters included the extraction well, two reinjection wells, two treatability study monitoring wells, and four existing monitoring wells. After the system was turned on, field parameters were measured weekly for all wells except for the extraction and reinjection wells, which were measured monthly. Two additional monitoring wells (62LF and 63LF) were installed in August 2000. After eight months of system operation, field parameters were measured biweekly. The standard field meters, instruments, and test kits used to make these measurements were calibrated in accordance with the manufacturer's instructions. Measurements were recorded on field data monitoring sheets.

6.3 Monthly Groundwater Sampling

Groundwater samples were collected prior to start-up of the treatability study to obtain baseline chemical and biochemical data in the study area. The samples were also collected monthly during the treatability study to track decreases and changes in chlorinated-solvent concentrations and daughter-product formation and destruction. Samples were also collected to help estimate the nutrient supplementation required during the study. Samples were analyzed for the contaminants/parameters listed in Table 6-2.

Table 6-2 Baseline Groundwater Sampling Protocol									
Analyte	Analytical Method	Wells Sampled	Purpose/Remarks						
VOCs	SW 8260	Extraction, injection, and TS monitoring wells	The purpose of VOC sampling is to obtain starting concentrations and track decreases in contaminant concentrations during the TS						
Metals 6010/7000		Extraction and TS monitoring wells (only newlyinstalled MW)	To examine clogging or solubilization effects on metals such as iron and manganese as a result of the created anaerobic-aerobic zone.						
Biological and Geochemica	l Parameters								
Total Kjeldahl Nitrogen	351.1 - 351.4	Extraction,	Nitrogen, phosphorus and carbon						
Ammonia-nitrogen	350.1	injection, and TS monitoring wells	measurements are required to estimate the amount and frequency of nutrient						
Total phosphorus	365.4	momentum would	supplementation required to optimize						
Orthophosphate	365.2 - 365.3		microbial activity. Chloride is a good indicator parameter used to estimate the						
Nitrate-nitrogen	352.1]	quantity of chlorinated solvents that have						
Total organic carbon	415.1]	been degraded during the TS.						
Chloride	325.3]							
Total heterotrophic counts	SM 9215B]							

Note:

TS = treatability study

Table 6-3 lists the protocol for periodic groundwater sampling and analysis. Groundwater samples were collected from area wells and analyzed for chemical and biochemical data.

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Table 6-3 Periodic Groundwater Sampling Protocol								
Analyte	Wells Sampled	Sampling Frequency						
VOCs	Extraction and TS monitoring wells	Monthly until the conclusion of the study						
Metals	Extraction and TS monitoring wells (only newly-installed MW)	After one month of treatability system operation						
Biological and Geochemical Paran	ieters							
Total Kjeldahl Nitrogen	Extraction, injection, and TS monitoring wells	Monthly until the conclusion of the						
Ammonia-nitrogen		study						
Total phosphorus								
Orthophosphate								
Nitrate-nitrogen								
Total organic carbon								
Chloride								
Total heterotrophic counts]							

Note:

TS = treatability study

6.4 Post-Shutdown Monitoring

After the system was shut down in December 2000, DO, pH, and ORP continued to be measured biweekly in samples from the wells to monitor changes in geochemistry in the aquifer. In March 2001, groundwater samples were collected from all the wells for VOC analysis. Post-shutdown monitoring was conducted to address the following:

- Will TCE concentrations rebound after the recirculation and amendment system is shutdown? If so, how long until concentrations increase?
- How long until the aquifer returns to pretreatment conditions?

• What happens to the *cis*-1,2-DCE that was generated from the reductive chlorination of TCE? Will persistent anaerobic condition result in *cis*-1,2-DCE degradation to VC? Conversely, is it possible that a return to aerobic conditions will foster *cis*-1,2-DCE degradation?

All sampling was performed in accordance with the Quality Assurance Project Plan (QAPP) and the Sampling and Analysis Program (SAP) developed as part of the RFI work plan for this site (E/A&H, 1994). Groundwater samples to be analyzed were sent to Laucks Testing Labs of Seattle, Washington, while groundwater samples for various gas analyses were sent to Microseeps of Pittsburgh, Pennsylvania.

7.0 RESULTS AND ANALYSIS

VOC (Section 7.1), geochemistry (Section 7.2), and hydraulic (Section 7.3) observations and

results are discussed below.

7.1 VOC Analysis

Select VOC analytical results for the treatability study monitoring wells, extraction well, and

reinjection wells are provided in Table 7-1. The table includes VOC data from the

baseline sampling event, which was performed prior to the start of the treatability study, from

nine subsequent events during the study (April to December), and from the post-shutdown event

(March 2001). Only the VOCs of prime concern in the pilot study area, PCE, TCE, cis-1,2-DCE,

and VC are included in this table. These data are also shown graphically in Appendix B.

Select phases of the A-A study are discussed below.

7.1.1 Early Concentration Fluctuations

Several significant TCE concentration fluctuations occurred during the first three months of

A-A system operation. The first sampling event (April) after system startup indicated

TCE concentrations in the extraction and two injection wells five to 30 times higher than

baseline (March) results. Comparatively, the TCE concentrations at the four initial downgradient

monitoring wells (007G58LFA, 007G58LFB, 007G59LFA, and 007G59LFB) were elevated in

March (210 to 2,100 μ g/L), substantially reduced in April (44 to 94 μ g/L), and elevated again in

May (190 to 1,200 μ g/L).

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Table 7-1 A-A Pilot Study Results (μg/L)												
		Baseline	e A-A Operation							Post- Shutdown		
Well ID	Parameter	3/00	4/00	5/00	6/00	7/00	8/00	9/00	10/00	11/00	12/00	3/01
007G03LF	PCE	47	41	51	66	91	110	140	130	150	120	73
upgradient monitoring	TCE	26	17	18	16	14	16	18	17	15	14	12
well	cis-1,2-DCE	<3	<3	<3	<3	<3	<3	<3	<3	<3	<3	<3
	VC	<3	<3	<3	<3	<3	<3	<3	<3	<3	<3	<3
007G57LF	PCE	7	9	45	97	70 J	68	52	44	39	38	45
extraction well	TCE	160	2,400 D	4,400 D	3,100 D	2,300	2,500 D	2,300 D	1,400 D	1,500 D	1,000 D	1,600 D
	cis-1,2-DCE	2 Ј	4	11	12	< 150	12	10	14	24	42	83
	VC	<3	<3	<3	<3	<150	<3	<3	<3	<3	<3	<3
007G60LF	PCE	50	7	30	53	66	54	48	NS	NS	28	<3
injection well	TCE	66	2,200 D	3,000 D	2,300 D	2,100	1,900 D	1,800 D	NS	NS	1,300 D	2 Ј
	cis-1,2-DCE	<3	4	7.7	8	<60	11	22	NS	NS	54	2 J
	VC	<3	<3	<3	<3	<60	<3	<3	NS	NS	<3	540 D
007G61LF injection well	PCE	41	6 J	35	50	75	55	47	NS	NS	30	<3
	TCE	300 D	1,600 D	3,400 D	2,300 D	2,200	1,900 D	1,800 D	NS	NS	1,400 D	<3
	cis-1,2-DCE	4	5 J	8.3	7	<60	11	17	NS	NS	57	<3
	VC	<3	<6	<3	<3	<60	<3	<3	NS	NS	<3	480 D

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Table 7-1 A-A Pilot Study Results (μg/L)												
		Baseline	ne A-A Operation									Post- Shutdown
Well ID	Parameter	3/00	4/00	5/00	6/00	7/00	8/00	9/00	10/00	11/00	12/00	3/01
007G62LFA	PCE	NA	NA	NA	NA	NA	22	14	10	6	<3	<3
intermediate monitoring	TCE	NA	NA	NA	NA	NA	700 D	360 D	250 D	150	2 J	22
well	cis-1,2-DCE	NA	NA	NA	NA	NA	940 D	990 D	1,100 D	1,100 D	1,100 D	990 D
	VC	NA	NA	NA	NA	NA	<3	<3	<3	<3	<3	81
007G62LFB	PCE	NA	NA	NA	NA	NA	28	16	9	7	<3	<3
intermediate monitoring	TCE	NA	NA	NA	NA	NA	820 D	460 D	270 D	190	5	<3
well	cis-1,2-DCE	NA	NA	NA	NA	NA	880 D	990 D	1,000 D	1,100 D	1,100 D	1,000 D
	VC	NA	NA	NA	NA	NA	<3	<3	<3	<3	<3	92
007G63LFA	PCE	NA	NA	NA	NA	NA	50	45	38	35	19	<3
intermediate monitoring	TCE	NA	NA	NA	NA	NA	1,800 D	1,600 D	1,300 D	1,200 D	530 D	10
well	cis-1,2-DCE	NA	NA	NA	NA	NA	22	140	110	270 D	560 D	1,000 D
	VC	NA	NA	NA	NA	NA	<3	<3	<3	<3	<3	<3
007G63LFB	PCE	NA	NA	NA	NA	NA	53	44	38	33	21	<3
intermediate monitoring	TCE	NA	NA	NA	NA	NA	1,700 D	1,600D	1,300 D	1,200 D	560 D	<3
well	cis-1,2-DCE	NA	NA	NA	NA	NA	43	150	140	240 D	520 D	1,200 D
	VC	NA	NA	NA	NA	NA	<3	<3	<3	<3	<3	4

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Table 7-1 A-A Pilot Study Results (μg/L)												
	Baseline A-A Operation									Post- Shutdown		
Well ID	Parameter	3/00	4/00	5/00	6/00	7/00	8/00	9/00	10/00	11/00	12/00	3/01
007G58LFA	PCE	9	1 J	4.8	19	55 J	53	41	22	15	10	34
intermediate monitoring	TCE	740 D	77	920 D	1,800 D	2700	2,300 D	1,700 D	800 D	710 D	370 D	1,300 D
well	cis-1,2-DCE	5	<3	2.1 J	4	< 60	11	<3	620 D	840 D	810 D	150 D
	VC	<3	<3	<3	<3	<60	<3	<3	<3	<3	<3	<3
007G58LFB	PCE	3	1 J	6.5	21	63	59	39	13	5	7	32
intermediate monitoring	TCE	210 D	69	1,200 D	2,000 D	2,600	2,500 D	1,500 D	400 D	140	170	1,300 D
well	cis-1,2-DCE	2 J	<3	2.7 Ј	5	<60	13	610 D	930 D	1,300 D	1,000 D	170
	VC	<3	<3	<3	<3	<60	<3	<3	<3	<3	<3	<3
007G59LFA	PCE	20	5	3.1	18	<60	35	49	34	24	12	29
intermediate monitoring	TCE	2,100 D	94	190 D	1,700 D	2,000	1,900 D	2,100 D	1,200 D	960 D	570 D	910 D
well	cis-1,2-DCE	18	1 J	1.1 J	7	< 60	9	110	210 D	450 D	720 D	320 D
	VC	<3	<3	<3	<3	< 60	<3	<3	<3	<3	<3	<3
007G59LFB	PCE	11	3 J	4.2	18	46 J	55	43	33	19	8	25
intermediate monitoring	TCE	1,600 D	44	560 D	1,800 D	2,400	2,500 D	1,700 D	1,100 D	850 D	370 D	920 D
well	cis-1,2-DCE	14	<3	2 J	6	< 60	11	230 D	360 D	620 D	880 D	450 D
	VC	<3	<3	<3	<3	<60	<3	<3	<3	<3	<3	1 J

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Table 7-1 A-A Pilot Study Results (μg/L)												
Baseline A-A Operation										Post- Shutdown		
Well ID	Parameter	3/00	4/00	5/00	6/00	7/00	8/00	9/00	10/00	11/00	12/00	3/01
007G04LF	PCE	17	110	5.2	10	35 J	38	49	35	27	<3	22
intermediate monitoring	TCE	2,000 D	1,800 D	240 D	1,200 D	2,400	2,400 D	2,300 D	1,400 D	1,100 D	3 J	930 D
well	cis-1,2-DCE	2 J	2 J	1 J	3 J	<60	12	36	260 D	540 D	<3	260 D
	VC	<3	<3	<3	<3	<60	<3	<3	<3	<3	<3	<3
007G04UF	PCE	<3	<3	<3	2 J	<3	<3	<3	9	4	15	<3
intermediate monitoring	TCE	2 J	1 J	190 D	590 D	15	130	74	400 D	160	800 D	1 J
well	cis-1,2-DCE	<3	<3	5.7	15	<3	3 J	2 J	8	4	580 D	<3
	VC	<3	<3	<3	<3	<3	<3	<3	<3	<3	<3	<3
007G21LF	PCE	6	6	6.4	5	7	6	8	NS	NS	9	12
downgradient monitoring	TCE	26	27	26	21	25	26	31	NS	NS	43	52
well	cis-1,2-DCE	<3	<3	<3	<3	<3	<3	<3	NS	NS	1 J	<3
	VC	<3	<3	<3	<3	<3	<3	<3	NS	NS	<3	<3

Notes:

A secondary dilution was used to analyze the sample. Estimated value. D

not applicable (wells not installed until August 2000). NA

not sampled. NS

The TCE concentration fluctuations are likely attributed to hydraulic effects from the startup of the recirculation system. Groundwater with relatively high TCE concentrations was drawn to the extraction well during system operation. This TCE-contaminated groundwater was eventually extracted and the re-injected into wells 007G60LF and 007G61LF, which resulted in the significant concentration increase at those locations. The downgradient monitoring wells also exhibited concentration changes because of the pumping-enhanced movement of TCE in the subsurface. Biological induced changes did not likely occur until later in the evaluation (August).

7.1.2 PCE and TCE Reduction and Formation of Daughter Products

The pumping-induced TCE concentration fluctuations seemingly stabilized by June. Injection well TCE concentrations were consistent with extraction well water quality. Though indicating hydraulic stabilization, downgradient well (007G58LF and 007G59LF) sampling results from June, July, and August also suggested minimal impact from the A-A system. In response, the carbon source was switched from fructose to acetate in June in an effort to accelerate the development of anaerobic conditions. When little impact was observed in July, two additional monitoring wells (007G62LF and 007G63LF) were installed closer to the injection wells to evaluate whether TCE reduction was actually occurring closer to the injection wells and had not yet impacted the wells that were further downgradient.

August sampling results from the new wells indicated elevated levels of *cis*-1,2-DCE (particularly in 007G62LF), which is a common biological daughter product of PCE and TCE reductive dechlorination. Because *cis*-1,2-DCE concentrations were very low before the evaluation was started, its significant increase in the aquifer indicates that the amendments (fructose/acetate and ammonium phosphate) effectively triggered reductive dechlorination. Moreover, *cis*-1,2-DCE concentrations remained low at background well 007G3LF and (considerably) downgradient well 21LF, further confirming that it was the augmentation that induced its formation. In addition, *cis*-1,2-DCE development during the study also shows that the fluvial aquifer does possess

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the required consortia of microorganisms to reduce PCE and TCE under the appropriate

geochemical conditions.

Because wells 007G62LF and 007G63LF were installed after the carbon source was changed from

fructose to acetate, it was difficult to evaluate whether fructose is an adequate amendment. If the

newest monitoring wells had been installed at the beginning of the study, it is possible that PCE

and TCE reduction would have been observed earlier and fructose would have remained the

carbon source of choice. Because of time constraints, carbon source selection and treatment time

were not evaluated independently.

From August through system shutdown in December, all four downgradient monitoring wells

(007G58LF, 007G59LF, 007G62LF, and 007G63LF) exhibited substantial PCE and

TCE concentration decreases and simultaneous cis-1,2-DCE increases. Though an order of

magnitude lower in concentration than TCE, PCE concentrations also discernibly deceased during

the study. Also, there did not appear to be any significant differences in concentrations between

groundwater in the upper and lower portions of the fluvial deposits.

Before reductive dechlorination occurred in the latter half of the evaluation, there was a period

from May through July in which carbon in the form of fructose and sodium acetate was depleting

the natural oxygen via microbial activity. Following oxygen depletion, the groundwater system

gradually developed the appropriate biochemical environment, which included a

microbial acclimation process to degrade chlorinated solvents.

While the formation of *cis*-1,2-DCE is an excellent indicator that the aquifer became anaerobic and

was able to sustain reductive dechlorination of PCE and TCE, the data also suggest that this

intermediate compound did not degrade as quickly as it was being formed under the

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treatability-study conditions. As the study progressed, cis-1,2-DCE began appearing

further downgradient in the study area as it was detected at significant concentrations at

well 007G04LF.

As such, the redox state of groundwater in the aquifer may not have been low enough for the

continued reductive dechlorination of cis-1,2-DCE to VC during the recirculation phase of the

study. Again, cis-1,2-DCE can also degrade aerobically or cometabolically, but because the study

area was devoid of dissolved oxygen, this pathway was not likely. Finally, it is also possible that

cis-1,2-DCE degradation requires the acclimation of a consortia of microorganisms different from

those that degrade PCE and TCE. The buildup of these microorganisms may include a lag period,

which was not established in the aquifer during the pilot study.

VC was not detected in the aquifer in the treatability area wells during the recirculation phase of

the study because it was either not being created or was degrading very quickly upon formation.

VC is known to degrade under aerobic and anaerobic conditions. For this particular application,

VC can be anaerobically oxidized by native microorganisms capable of using ferrous iron as the

electron acceptor. In anaerobic aquifers with relatively high concentrations of natural iron

(>1 mg/L) (see Table 7-2), VC could degrade upon being formed via reductive dechlorination.

Though it is conceivable that VC was being degraded upon formation via iron-mediated metabolic

activity, it is more likely that no VC was ever formed during the recirculation phase of the study.

Table 7-2 Iron Concentrations (μg/L)								
Sample Date								
Well	Well 3/6/00 4/12/00							
007G57LF	5,020	3,220						
007G58LFA	1,570	1,620						
007G58LFB	3,260	1,640						

7.1.3 Post-Shutdown Monitoring

After the system was shut down in December 2000, DO, pH, and ORP continued to be measured biweekly in samples from the wells to monitor changes in geochemistry in the aquifer. In March 2001, groundwater samples were collected from all the wells for VOC analysis. As discussed in Section 6.4, post-shutdown was conducted to address TCE rebounding, the fate of *cis*-1,2-DCE, and aquifer re-aeration.

There were several interesting results three months after the extraction-re-injection system was stopped. Post-shutdown results for the extraction well indicated increased TCE and cis-1,2-DCE concentrations. The apparent TCE increase could be attributed to an anomalously low TCE concentration December (October, November, December, and March data were 1,400, 1,500, I,000, and 1,600 μ g/L respectively). However, the cis-1,2-DCE increase is consistent with its gradual increase since August. Without a concomitant TCE decrease at this well, this could be evidence that cis-1,2-DCE is forming upgradient of and migrating to the extraction well.

Stopping the recirculation system and slowly returning the aquifer to natural hydraulic conditions resulted in a relatively stagnant environment, particularly near the injection wells (60LF and 007G61LF) and first row of monitoring wells (007G62LF and 007G63LF). This stagnancy

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coupled with ample1 residual organic carbon in the groundwater resulted in much

stronger anaerobic conditions resulting in cis-1,2-DCE degradation to VC (up to 540 μ g/L in

007G60LF). Consistent with the analytical results, ORP measurements taken from the injection

wells have ranged from -150 to -200 mV since system shutdown, likely low enough for

cis-1,2-DCE and VC degradation. Moving from the injection wells to the extraction well in the

direction of groundwater flow, ORP measurements are increasingly more positive and the quantity

of contaminant mass increases.

Though speculative as of March 2001, continued post-shutdown monitoring would be conducted

to further assess the following:

1. Sufficiently anaerobic conditions can be generated in the lower fluvial aquifer to promote

complete chlorinated VOC mineralization. If not, a return to natural aerobic conditions

would likely stimulate the degradation of cis-1,2-DCE and VC.

2. Indigenous microbes capable of degrading *cis*-1,2-DCE and VC are present.

3. With some TCE mass now reduced to *cis*-1,2-DCE or VC, further degradation will likely

occur in either sustained anaerobic or naturally re-aerated pre-treatment conditions.

Continued monitoring will demonstrate whether intrinsic aerobic conditions can sustain

biodegradation or sparging/natural aerobic conditions are required to complete the

treatment process.

¹ Compared with pre-shutdown samples, TOC concentrations in samples from the injection wells and 007G62LF and 007G63LF monitoring wells were one to two orders of magnitude higher after the system was

shut down.

7.1.4 Degradation Rates

Microbial kinetics are generally dependent on several factors:

- microbial growth and populations
- starting concentrations
- availability of carbon sources and nutrients,
- degradation occurring due to attached-phase and suspended-phase microorganisms.

The zero-order degradation rate equation used to calculate contaminant degradation rates is:

$$C_t = C_0$$
-kt

Where:

 C_t = Solute (TCE) concentration in mg/L at time t

 C_0 = Solute (TCE) concentration in mg/L at time "zero" (in this case, August)

t = time (days)

 $k = first-order degradation rate in day^{-1} (1/day)$

Comparatively, the first-order degradation rate equation, which commonly represents TCE degradation kinetics, used to calculate contaminant degradation rates is:

$$C_t = C_0 e^{-kt}$$

PCE and TCE degradation rates for the purpose of estimating clean-up times and scale-up factors were estimated for the two closest monitoring wells (007G62LF and 007G63LF) using August, September, October, November, and March VOC results. The December data were considered anomalous and not included the assessment.

The data were plotted arithmetically and logarithmically for each of the wells to estimate the overall zero- or first-order degradation rate, which accounts for both biological and physical (i.e., dispersion, adsorption, volatilization, and dilution) degradation mechanisms. The graphs for each of the wells are provided in Appendix C. The results of the degradation rate assessment are summarized in Table 7-3.

	Table 7-3 Degradation Rate Estimation Summary											
	•				Date/Day							
			8/22/00	9/21/00	10/18/00	11/14/00	3/27/01					
Well	voc	Туре	0	30	57	84	217	\mathbb{R}^2	k	0		
007G62LFA	TCE	Meas.	700 D	360 D	250 D	150	22	0.995	-0.016	1		
,		Model	700	439	288	189	24	0.993	-0.010	1		
	PCE	Meas.	22	14	10	6	<3	0.921	-0.009			
		Model	22	17	13	10	<3	0.921	-0.009	1		
007G62LFB	TCE	Meas.	820 D	460 D	270 D	190	<3	0.983	-0.026	1		
		Model	820	373	183	90	<3	0.963	-0.020	1		
	PCE	Meas.	28	16	9	7	<3	0.905	-0.010	1		
		Model	28	21	16	13	<3	0.903	-0.010	1		
007G63LFA	TCE	Meas.	1,800 D	1,600 D	1,300 D	1,200 D	10	0.995	-8.286	0		
. •		Model	1800	1551	1328	1104	2J	0.993	-0.200	U		
	PCE	Meas.	50	45	38	35	<3	0.995	-0.219	0		
		Model	50	43	<i>3</i> 8	32	<3	0.993	-0.219			
007G63LFB	TCE	Meas.	1,700 D	1,600 D	1,300 D	1,200 D	3	0.989	9 045	0		
		Model	1700	1459	1241	1024	-46	0.969	-8.045	0		
	PCE	Meas.	53	44	38	33	3	0.998	-0.226	0		
		Model	53	46	40	34	4	0.998	-0.226	U		

Notes:

D = A secondary dilution was used to analyze the sample.

J = Estimated value. $R^2 = R$ -squared value

k = degradation rate constant (1/day)

O = order of the reaction

Interestingly, PCE and TCE degradation rates for well 007G62LF are represented by first-order kinetics while 007G63LF is represented by zero-order kinetics. The difference between the two wells is likely due to starting concentrations. PCE and TCE concentrations at well 007G63LF are approximately twice those at 007G62LF.

PCE

Zero-order rates for PCE ranged from 0.219 day⁻¹ (80 yr⁻¹) to 0.226 day⁻¹ (83 yr⁻¹). First-order rates for PCE ranged from 0.009 day⁻¹ (3.3 yr⁻¹) to 0.010 day⁻¹ (3.7 yr⁻¹). As shown in Table 7-4, first-order PCE degradation rates from the A-A pilot study are reasonably consistent with other enhanced bioremediation evaluations.

	Table 7-4 PCE Degradation Rate Summary										
	De										
Source	Day ⁻¹	Year ⁻¹	Half-Life (days)	Comments							
A-A Study	0.009 - 0.01	3.3 - 3.7	69 - 77								
Sheldon, 1999	0.005 - 0.011	2.0 - 3.8	66 - 128	rates decreased as HRC was depleted							
Dooley, 1999	0.021	7.7	33	rate order not reported (assumed first)							
Maierle, 2001	0.021 - 0.027	7.7 - 9.5	25.7 - 33	enhanced reductive dechlorination (ERD)							

TCE

Zero-order rates for TCE ranged from 8.05 day⁻¹ (2,940 yr⁻¹) to 8.29 day⁻¹ (3,025 yr⁻¹). First-order rates for TCE ranged from 0.016 day⁻¹ (5.8 yr⁻¹) to 0.026 day⁻¹ (9.5 yr⁻¹). As shown in Table 7-5, first-order TCE degradation rates from the A-A pilot study are reasonably consistent with other enhanced bioremediation evaluations. Further, the higher estimated first-order value

is about an order of magnitude greater than the reported literature value of natural attenuation studies at monitored natural attenuation (MNA) sites; the lower estimated value is five times higher. The degradation rate comparison indicates that the site is very amenable to microbial enhancement.

	Table 7-5 TCE Degradation Rate Summary											
	De	gradation Ra	tes									
Source	Day ⁻¹	Year ⁻¹	Half-Life (days)	Comments								
Enhanced In Situ Bioremediation												
A-A Study	0.016 - 0.026	5.8 - 9.5	26.7 - 43.3									
Dooley, 1999	0.018	6.6	38.5	rate order not reported								
Maierle, 2001	0.005 - 0.023	1.8 - 8.4	30.1 - 138.6	enhanced reductive dechlorination (ERD)								
MNA												
Cox, 1995	0.003	1.1	231	sequential anaerobic-aerobic aquifer								
Lee, 1995	0.002	0.7	347	downgradient of an industrial landfill								

7.1.5 TCE Mass Degradation Estimation

This section is a preliminary effort at estimating the mass of TCE that was degraded as a result of the treatability study. Contours were drawn to estimate the affected area using TCE concentrations in the fluvial deposit wells in the study area. Rather than relying on the data from the beginning and end of the study because of pumping-induced fluctuations, concentration contours were generated for all 11 sampling events (baseline, nine months of operation, and one shut-down event) to evaluate TCE mass changes over time. The concentration contours for each month are shown in Figure 7-1.

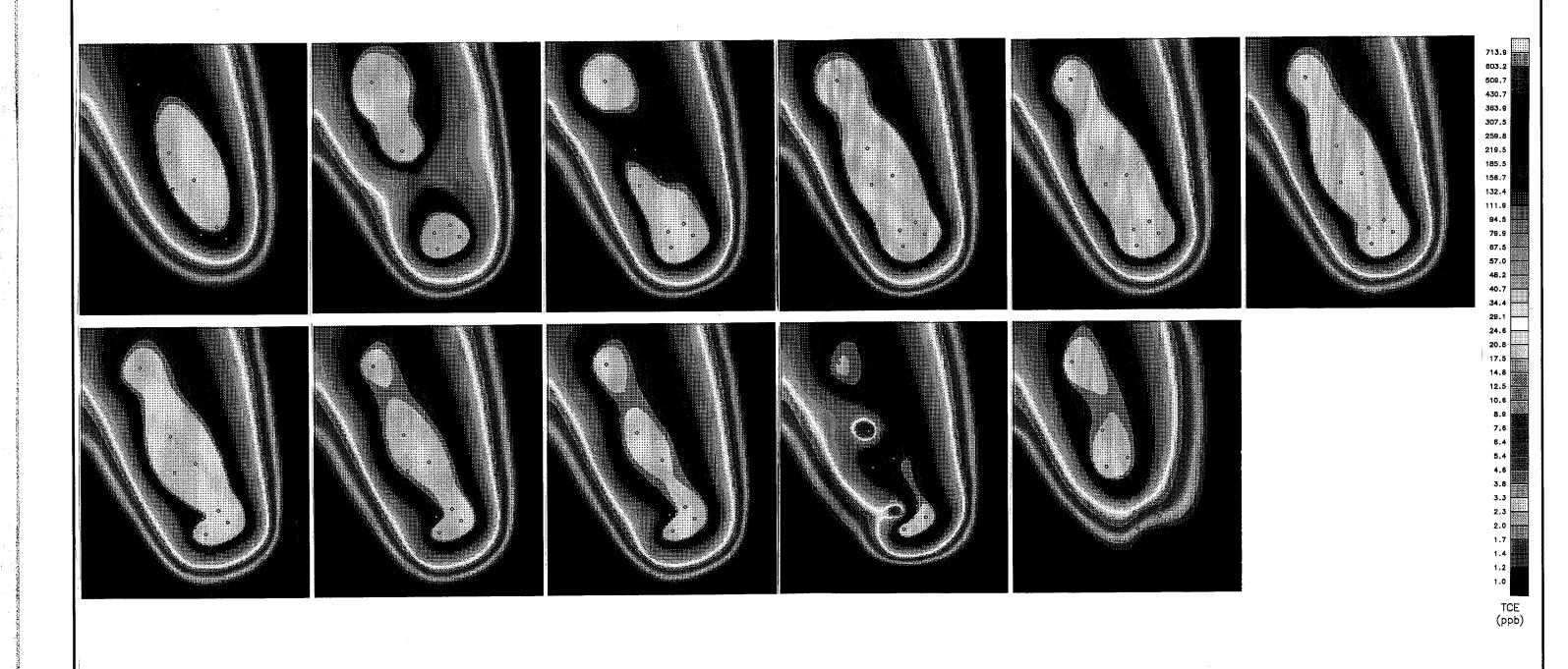




FIGURE 7-1
MONTHLY TCE PLUMES
A-A SEQUENTIAL STUDY TREATABILITY REPORT
AOC A - NORTHSIDE FLUVIAL GROUNDWATER
NSA MID-SOUTH, MILLINGTON, TN

Then, for simplicity, the 5, 100, and 1,000 μ g/L contours were used to generate TCE mass estimates for each month during the study. The area within each of the three contour intervals was multiplied by an assumed saturated thickness (40 feet), porosity (25%), and average concentration within the contour to calculate the mass of TCE for each month. The results are summarized in Table 7-6. TCE mass trends are presented graphically in Figures 7-2a and b.

	•			TCI		le 7-6 mation Sum	mary			•	
ppb	Mar-00	Apr-00	May-00	Jun-00	Jul-00	Aug-00	Sep-00	Oct-00	Nov-00	Dec-00	Mar-01
	TCE Plume Area (square feet)										
5	16,777	18,960	18,548	18,666	18,038	18,146	18,777	18,298	18,690	17,612	15,066
100	9,805	8,644	11,608	12,050	11,372	11,457	11,703	11,280	11,289	9,161	8,120
1,000	2,274	1,857	1,940	4,062	3,727	3,516	3,067	1,109	660	169	584
! !	TCE Mass (kg)										
5	0.69	0.78	0.76	0.77	0.74	0.75	0.77	0.75	0.77	0.73	0.62
100	4.04	3.56	4.78	4.96	4.68	4.72	4.82	4.64	4.65	3.77	3.34
1,000	6.24	5.10	5.33	11.15	10.23	9.65	8.42	3.04	1.81	0.46	1.60
Total	10.97	9.44	10.87	16.88	15.65	15.12	14.01	8.44	7.23	4.96	5.57
				Percent M	ass Change	from Baseli	ine (Mar-00))			
5	100%	113%	111%	111%	108%	108%	112%	109%	111%	105%	90%
100	100%	88%	118%	123%	116%	117%	119%	115%	115%	93%	83%
1000	100%	82%	85%	179%	164%	155%	135%	49%	29%	7%	26%
Total	100%	86%	99%	154%	143%	138%	128%	77%	66%	45%	51%

Assumptions

Aguifer thickness = 40 feet

Average concentration in 5 ppb contour = 30 ppb

Average concentration in 100 ppb contour = 300 ppb

Average concentration in 100 ppb contour = 2,000 ppb

Porosity = 0.25

Figure 7-2a AOC A Pilot Study TCE Plume Area Trends

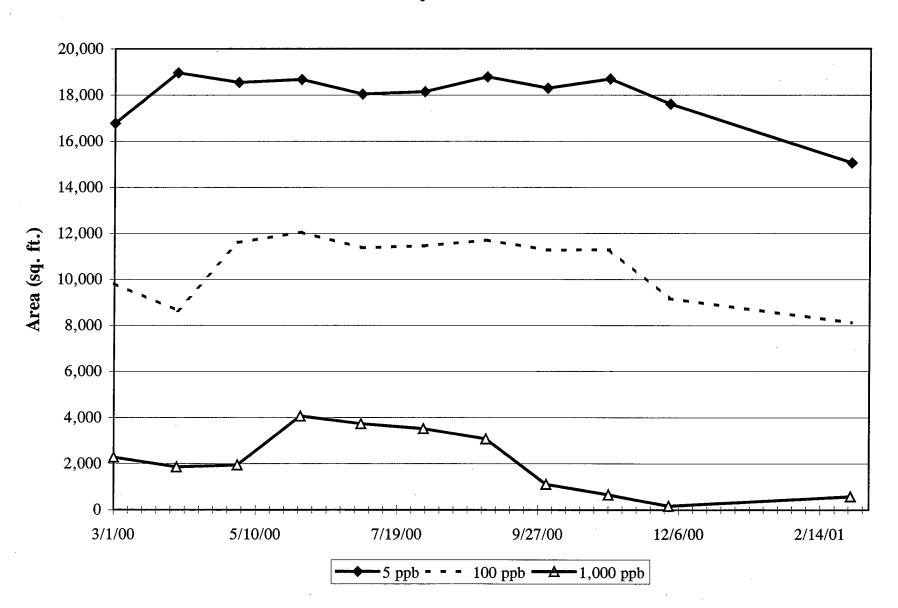
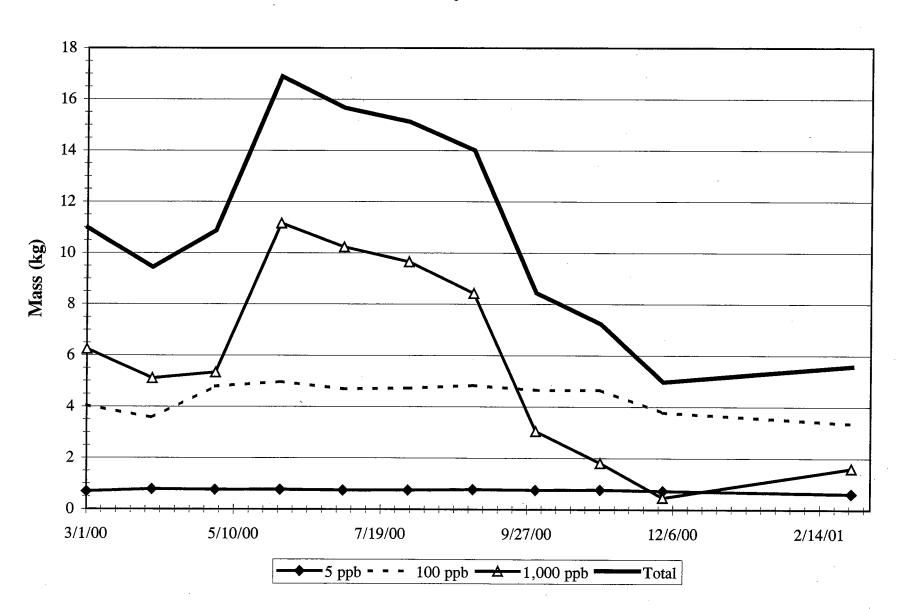


Figure 7-2b AOC A Pilot Study TCE Mass Trends



Discussion

As expected, the most significant mass change was observed in the 1,000 μ g/L plume. Relative to March 2000 (baseline sampling event), the TCE mass in this part of the plume decreased by approximately 75%. Coupled with the relatively minor decreases in the other portions of the pilot study area plume, total TCE mass decreased by 50% in one year (\approx 5 pounds). Comparatively, if measured from June 2000, when the 1,000 μ g/L plume was largest, the overall TCE mass actually decreased by 2/3 or about 10 pounds in nine months.

As shown on Figure 7-2a and b, and as discussed in Section 7.1.1, concentration and plume area fluctuations occurred during the pilot study. Initial fluctuations are likely attributed to the (1) hydraulic effects from the startup of the recirculation system as TCE was drawn towards the pilot study area, (2) microorganism acclimation delay, and (3) subtle variations in the contouring effort, which was done manually. Concentration, mass, and area changes during the latter portion of the study are a result of reductive dechlorination. Because the high concentrations were targeted by the injection system, and the microbes invariably acclimate faster to areas with higher electron donor and acceptor concentrations, the 1,000 μ g/L zone decreased first (August/September) followed by the 100 μ g/L (November) and the 5 μ g/L plume (November/ December).

7.2 Geochemistry Analysis

7.2.1 Field Data

When possible during the study, the field data were measured on a weekly or biweekly basis from the treatability monitoring wells and monthly from the extraction and reinjection wells. The results are summarized in Appendix D. The parameters include ammonia, CO₂, DO, ORP, and orthophosphate data. ORP data are also plotted on the VOC data graphs in Appendix B to demonstrate the relation between reducing conditions and reductive dechlorination of TCE and its daughter products.

DO and ORP

An overall observation of DO data indicates anaerobic conditions in the aquifer. Groundwater near the reinjection wells had very low DO concentrations from the middle of the study through system shutdown. Low ORP values were coincident with the low DO concentrations suggesting reducing conditions in the aquifer. Comparatively, DO concentrations were relatively elevated and ORP values were generally positive throughout the remainder of the study area until the system was shutdown. As soon as groundwater recirculation ceased, ORP values in the reinjection wells, both rows of downgradient monitoring wells, and 007G4LF became negative; some measurements were as low as -244 mV, which is sufficiently anaerobic for *cis*-1,2-DCE degradation.

Carbon Dioxide

CO₂ readings in Appendix D show that the critical monitoring wells 007G58LF, 007G59LF, 007G62LF, and 007G63LF did not show a discernible trend of any kind for this constituent. Moreover, CO₂ values in background well 007G3LF were in the same range as reported for the treatability areas wells. Therefore, for this treatability study, CO₂ readings may not be a valuable indicator of enhanced microbial activity.

7.2.2 Laboratory Data

Nutrient and geochemical data were collected monthly from the pilot study area wells and analyzed in the laboratory. Nutrient measurements include ammonia, phosphate, and TKN, while geochemical parameters of significance include TOC, nitrate, and chloride. These analyses were performed to check the adequacy of nitrogen and phosphorus as microbial nutrients in groundwater. The results are summarized in Appendix D.

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TOC:TKN:Phosphorus (C:N:P) Ratios

Ratios of TOC to TKN to phosphorus are summarized in Table 7-7. For most events and all sampled wells, these ratios show that nitrogen and phosphorus (the two most essential nutrients for microbial activity) are unlikely to have limited microbial activity in the aquifer. Based on an average composition of cell tissue of $C_5H_7NO_2$, about 12.4% by weight of nitrogen will be required. The phosphorous value is assumed to be one-fifth of this value (this equates to a 100:20:4 C:N:P ratio). These are typical values, not fixed quantities, because it has been shown that the percentage distribution of nitrogen and phosphorus in cell tissue varies with the age of the cell and environmental conditions (Metcalf & Eddy, 1991). For measurable events (i.e., when

TOC was detectable), the pilot-study nutrient ratios commonly met literature requirements. In

fact, as shown in Appendix D, nitrogen and phosphorus were above detection limits in some of

the samples in which TOC was not detected. As such, nitrogen and phosphorus were available

at adequate levels to sustain microbial degradation in the aquifer.

Iron

Iron was measured in three wells during the March and April sampling events. Results presented in Section 7.2.2 show significant concentrations of iron in the fluvial deposits that could assist the microbially-mediated reduction of VC under the appropriate conditions. This may be a critical factor when a mainly aerobic environment such as the fluvial deposits is converted to one that is more anaerobic. Under these converted conditions, the aerobic degradation of VC could be impeded and this daughter product could accumulate unless significant amounts of iron

could sustain degradation even under more reducing conditions.

Hydrogen Measurements

In May 2000, three wells were sampled for hydrogen and methane. Table 7-8 shows the results from this event. The low methane concentrations indicate that the aquifer was unlikely to be

methanogenic. The hydrogen values, however, indicate conditions in the aquifer were favorable

		· · · · · · · · · · · · · · · · · · ·			Table 7-7 C:N:P Ratios				-	
	Baseline					A-A Operation	1			
Well	3/00	4/00	5/00	6/00_	7/00	8/00	9/00	10/00	11/00	12/00
007G3LF	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	NS	NS	100:50:4
007G57LF	100:19:1.7	no carbon	100:33:10	no carbon	no carbon	no carbon	no carbon	100:ND:2.3	no carbon	100:ND:ND
007G60LF	100:35::4.1	no carbon	no carbon	no carbon	no carbon	no carbon	100:26:3.5	NS	NS	100:1.4:1.6
007G61LF	100:13:1.5	100:12:10	no carbon	no carbon	no carbon	100:18:4.2	100:2.4:0.9	NS	NS	100:1.2:ND
007G62LFA	NS	NS	NS	NS	NS	100:9.2:0.8	100:6.4:0.2	100:ND:0.5	100:ND:ND	100:ND:0
007G62LFB	NS	NS	NS	NS	NS	100:5.4:0.4	100:10:0.3	100:ND:0.5	100:ND:ND	100:0.3:0
007G63LFA	NS	NS	NS	NS	NS	100:4.4:0.3	100:13:0.5	100:ND:2.4	100:0.6:ND	100:0.2:0
007G63LFB	NS	NS	NS	NS	NS	100:4.5:0.8	100:11:0.9	100:ND:1.1	100:0.6:ND	100:0.20
007G58LFA	no carbon	no carbon	no carbon	no carbon	no carbon	100:ND:16	no carbon	no carbon	100:ND:ND	100:ND:10
007G58LFB	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	100:ND:ND	100:ND:286
007G59LFA	100:12:1.2	100:40:3	100:50:1.9	no carbon	no carbon	100:43:5.7	no carbon	no carbon	100:ND:ND	no carbon
007G59LFB	100:ND:2.7	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	100:ND:19	100:ND:ND	no carbon
007G4LF	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	100:ND:ND	100:ND:2.3
007G4UF	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	no carbon	100:ND:7.3	100:ND:ND	100:ND:1.1
007G21LF	no carbon	no carbon_	no carbon	no carbon	no carbon	100:60:4.5	no carbon	no carbon	NS:ND:ND	no carbon

Notes:

ND = nondetect

no carbon = TOC was below detection limits; therefore, no ratio was calculated

NS = not sampled

Table 7-8 Hydrogen and Methane Results (5/12/00)										
Well	Well Hydrogen (nM/L) Methane (μg/L)									
007G04LF	1.96	0.042								
007G04UF	2.02	0.050								
007G58LFA	2.07	0.038								
007G58LFB	1.65	0.043								

Note:

nM/L = nanomoles per liter

for sulfate-reduction and fairly conducive to reductive dechlorination of PCE and TCE. According to the *Technical Protocol for Evaluating Natural Attenuation of Chlorinated Solvents in Ground Water* (USEPA, 1998), hydrogen concentrations > 1 nM/L indicates the strong potential for reductive dechlorination. Further, if hydrogen concentrations are high enough to support sulfate reduction or methanogenesis, then reductive dechlorination is probably occurring, even if other geochemical indicators do not indicate that reductive dechlorination is possible (USEPA, 1998).

Heterotrophic Plate Count (HPC)

To monitor the microbial activity in the fluvial deposits aquifer, the wells were sampled monthly for HPC. The results for HPC are presented in Appendix D with the field and laboratory geochemical data. The mean values for all the wells ranged from 2.9×10^3 to 1.4×10^6 colony forming units (CFUs) per mL. There were no significant changes in any of the wells throughout the study. Most wells showed mean concentrations of 10^3 to 10^4 CFUs per mL. Aquifer augmentation did not appear to induce a significant increase in heterotrophic microbial populations. However, based on the *cis*-1,2-DCE production during the study, existing microorganisms appear to possess the capability of reducing TCE given the appropriate geochemical environments.

7.2.3 Post-Shutdown Geochemistry

Post-shutdown DO and ORP data indicate the continuation of reducing or anaerobic conditions three months after system shutdown. In particular, the aquifer in the vicinity of the two reinjection wells indicates the development of a reducing "stagnant" zone which is reflected in the sharp decrease in TCE and the buildup of VC. ORP readings in the direction of groundwater flow indicate less reducing conditions towards wells 007G62LF and 007G63LF and a gradual tendency towards the aerobic range at wells 007G58LF, 007G59LF, and 007G04LF, though groundwater still remains reducing in the entire treatability-study area. Over time, groundwater could become more aerobic and return to post-treatability conditions as the added carbon (acetate and fructose) is completely consumed.

7.3 Hydraulic Analysis

Based on groundwater elevations measured during the RFI, groundwater flows primarily to the northwest with localized small scale deviations. The hydraulic performance of the sequential A-A system was evaluated through (1) qualitative analysis of the potentiometric surface as measured during sequential sampling, and (2) analytical flow analysis of the potentiometric surface as measured during the sequential sampling.

7.3.1 Qualitative Analysis

Appendix E presents the water levels measured in each of the monitoring wells during the study, and Figures 1 through 8 in Appendix F present the modeled piezometric surface for each measurement event. The potentiometric contours are generated by statistical means only and do not take into account other factors (e.g., geology).

The measurement events included a baseline (3/7/00), a system start up (3/14/00), a series of regular monthly events (3/23/00 to 11/15/00), and a shutdown (12/12/00). Because the emphasis for this analysis is on the performance of the treatment system, only events that had

water-level measurements for the extraction and re-injection wells were used in the analyses (baseline, start-up, and six sampling events covering a 180-day period of performance).

The baseline condition potentiometric surface exhibits a "tongued" high in the general area of monitoring wells 007G04LF and 007G04UF, with a longitudinal axis directed in a northwesterly azimuth (Figure 1, Appendix F). Northeast of this the gradient azimuth and magnitude is relatively constant towards the north-northwest. Following system startup, the potentiometric surfaces from 4/10/00 to 7/16/00 also exhibit a similar extension downgradient, but the axis deviates somewhat in direction and magnitude. This may be associated with the injection of groundwater, which could create a downgradient extension of higher elevation isopleths. The direction and magnitude would vary depending on the injection rate and the extraction rate from well 007G57LF. The potentiometric surface for 8/21/00 exhibits a retraction of lower elevation ispoleths towards the extraction well, with an axis azimuth towards the southwest. This is related to a relative rise in the head in injection well 007G60LF, which was the results of screen clogging from fine-grained sediment and high levels of iron hydroxides.

After this date, a filter system was installed to help prevent recurrence of this relative rise in water level. The 9/19/00 potentiometric surface, which is the final surface profiled during the 180-day period of performance, exhibits a high near the extraction well, and a subdued "tonguing" of the higher area towards the northwest, which is what would be expected during a static operation of the extraction/injection hydraulics. Unfortunately, the surface also exhibits a relative low to the southeast of well 007G60LF; this is a statistical extrapolation of the lower elevation of this well versus that of 007G61LF, 007G62LF and 007G63LF, and should not be considered as "real." The only alternative to eliminate this feature for the particle tracking (see Section 7.3.2) is to insert "dummy points," an exercise which was not conducted in the interest

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of presenting only valid data. In summary, conclusions that may be inferred from this evaluation include:

• Although some local rise in the potentiometric surface was noted in the vicinity of the monitoring wells between the injection and extraction wells as a baseline condition, the observed rise may be the result of drilling artifacts.

Operation of the system appears to have been quite variable over the performance period, resulting in a number of perturbations of the basic potentiometric configuration. However, by 9/19/00, hydraulic conditions appear to be consistent with the intended design for injection and extraction.

7.3.2 Quantitative Analysis

To assess the hydraulic performance of the system, particle tracking modeling was conducted for the period of operation. This entailed entering the grid file for each potentiometric surface, along with appropriate aquifer parameter information, into GWPath (Schafer, 1992), a commercially available software platform for forward and reverse pathline analysis. The reader is referred to the treatability study work plan for specifics on GWPath. Table 7-9 provides the input parameters used for the analyses. For each of the sequential sampling events from April through September, reverse tracking was conducted for the extraction well, and forward tracking was conducted for each of the re-injection wells. Figures 9 through 14 in Appendix F present the results of these exercises. Similar to the qualitative evaluation, only events that included extraction and re-injection well measurements (post system start-up) were included for analysis.

Pathlines illustrating zones of influence for the period of the first 60 days exhibit very little movement of groundwater towards the hydraulic sources and sinks. The period of the next 60 days exhibits a fairly consistent movement of water away from the injection wells, but

Table 7-9 GWPath Input Parameters									
Parameter	Value	Source/Logic							
Flow domain	Horizontal	Lateral pathline analysis.							
Model domain grid	$X_{min} = 813673; X_{max} = 813897$ $Y_{min} = 391936; Y_{max} = 392560$ $X_{spacing} = 6.4; Y_{spacing} = 13$ Rows = 36; Columns = 49	Incorporation of all wells monitored.							
Hydraulic conductivity	X = 5 ft/d; Y = 5 ft/d	Design memo; Assumed lateral isotropy.							
Porosity	25%	Value used by USGS in transport modeling; from core sample.							
Hydraulic head file	Grid file for each sampling event pot surface	Based on model domain grid specs.							
Travel time	Determined for each sampling event; 3/14/00 as start time.	Period of operation from start-up.							
Output	Extraction well — reverse tracking; Re-injection well — forward tracking	Lateral tracking of pathlines.							

very little movement towards the extraction well. This is a function of the relatively low gradient mapped in the vicinity of well 007G57LF. The pathline analysis for the 8-21-00 period, which had a considerably different potentiometric configuration, demonstrates the effect of the steeper hydraulic gradient from the injection wells towards the northeast. Finally, the pathline analysis for the period 9-19-00 shows the response of pathlines towards the hydraulic sink of well 007G60LF.

Theoretically, the forward pathlines from the injection wells would extend to the northwest towards the extraction well. Similar to the other periods, however, the extraction well influence appears to be very small, which likely is a result of the flat gradient in the vicinity of 007G57LF.

Pathline azimuths are not identical for each evaluation period as they are influenced by the selected potentiometric surface. The azimuths reflect the gradient direction, which was noted to

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change, likely as a function of the variable injection rates. Therefore, the system, through the

period of performance, operated as a highly transient system. The optimal static conditions (that

is, relative highs near the injection wells, with a capture influence associated with the

lower downgradient extraction well) may require more than the 180 days of operation to establish.

In summary, the hydraulic performance analytical evaluation indicates that injected water did not

migrate into the capture zone of the extraction well during the limited time period of the

pilot study, nor did the migration of injected water stay consistent in terms of direction.

Extracted water, based on pathline analysis only, is derived from the immediate vicinity of the

extraction well, which is thought to be a function of the relatively flat gradient.

Water-Level Trends

Over the course of the study, water levels generally exhibited an increase through May, and then

a general decrease until December. To evaluate the uniformity or non-uniformity of these trends,

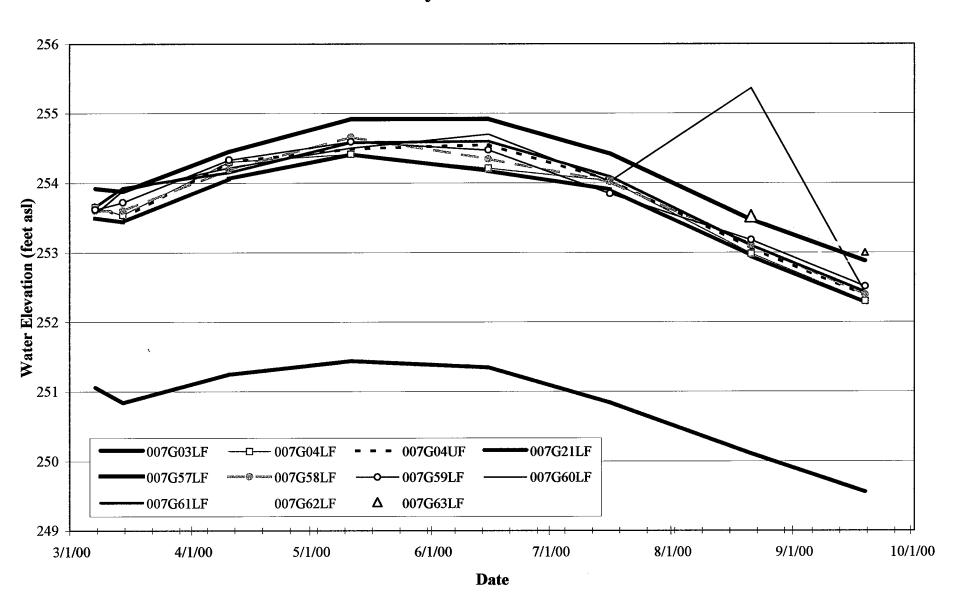
water-level trends for each well were plotted, and are presented as Figure 7-3. As demonstrated

by the graph, all wells demonstrate similar trends, and thus were somewhat equally affected, with

the exception of an increase in head in well 007G60LF during August. This exception may be a

reflection of an increase in injection rate in that particular well in July.

Figure 7-3
A-A System Water Levels



8.0 CONCLUSIONS AND RECOMMENDATIONS

Conclusions

The main objective of performing the A-A treatability study in the fluvial deposits aquifer at the selected location within AOC A was to determine the feasibility and effectiveness of using enhanced in situ bioremediation to treat PCE- and TCE-contaminated groundwater. Historically, the fluvial deposits aquifer in the treatability area has been slightly aerobic and has very low concentrations of natural carbon. Therefore, nutrients (fructose/acetate and ammonium phosphate) were added during the study to stimulate indigenous microorganisms to change the redox state of the aquifer.

After the nine-month treatability study, it appeared that reductive dechlorination of PCE and TCE is feasible via bioaugmentation. The attainment of reducing conditions was confirmed by negative ORP measurements, low DO concentrations, and elevated hydrogen concentrations during field geochemical sampling. However, the most significant observation was the two-order-magnitude increase in *cis*-1,2-DCE concentrations in the study area monitoring wells.

Pilot study results also indicated that *cis*-1,2-DCE did not degrade at a rate commensurate to its formation during system operation. It accumulated because of persistent anaerobic conditions where natural aerobic conditions were expected to stimulate its degradation. *Cis*-1,2-DCE concentrations increased over time and were measured farther downgradient as the study progressed. As such, sparging would likely be required in a full-scale system to accelerate its removal.

Preliminary TCE mass balances and parent-compound degradation rates were also performed during the evaluation. Preliminary estimates indicate a 50% reduction in TCE mass in the area. Zero-order rates for PCE ranged from 0.219 day⁻¹ (80 yr⁻¹) to 0.226 day⁻¹ (83 yr⁻¹) while first-order rates ranged from 0.009 day⁻¹ (3.3 yr⁻¹) to 0.010 day⁻¹ (3.7 yr⁻¹). Comparatively,

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Section 8: Conclusions and Recommendations

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zero-order rates for TCE ranged from 8.05 day⁻¹ (2,940 yr⁻¹) to 8.29 day⁻¹ (3,025 yr⁻¹) and first-order rates ranged from 0.016 day⁻¹ (5.8 yr⁻¹) to 0.026 day⁻¹ (9.5 yr⁻¹). These numbers were compared to literature values for natural attenuation sites. The comparison shows rates of removal of TCE equal to or greater than reported values.

Although the system operation appeared to be variable over the study period, hydraulic conditions appeared to be consistent with the extraction and injection system design by about six months after start up. The analytical evaluation of the hydraulic performance indicates that injected water has not yet migrated to the capture zone of the extraction well during the limited period of operation of the pilot study, nor has the migration of injected water been consistent in terms of direction. The extracted water was shown to have originated from the immediate vicinity of the extraction well, which is a function of the relatively flat hydraulic gradient.

Chemical and geochemical data from the treatability-study wells collected three months after system shutdown indicate that the system was gradually returning to pre-treatability conditions. VC was detected at the reinjection wells for the first time as a result of the creation of a "stagnant" reducing zone in the vicinity of the wells. VC, as well as *cis*-DCE are expected to gradually decrease over time along the groundwater flow path. Overall, the groundwater in the vicinity of the injection wells remains anaerobic but is likely to gradually turn aerobic after all the remaining augmented carbon is consumed.

Recommendations

The A-A treatability study was the first attempt to provide an engineered solution to remedying the highest concentrations of chlorinated solvent contamination in the aquifer fluvial deposits at AOC A. The results show that this technology can feasiblely reduce TCE at AOC A. Preliminary calculations of degradation rates and mass reductions should be used during the CMS and reported as a baseline against which other treatment alternatives can be compared.

A-A Sequential Remediation Treatability Study Report AOC A — NSA Mid-South

Section 8: Conclusions and Recommendations

Revision 1; May 17, 2002

Preliminary calculations of degradation rates and mass reductions should be used during the CMS

and reported as a baseline against which other treatment alternatives can be compared.

The location, spacing, and number of extraction, reinjection, and pumping wells for the

treatibility study were based on site investigation data and hydrogeological modeling. This

information can be used to design a treatment system at other locations at AOC A where treatment

is required.

This study used two carbon sources, fructose and sodium acetate. Fructose or molasses is the

preferred carbon source because cost of carbon amendment would be a significant factor in

deciding the type of bioaugmentation for treatment of larger areas or areas with higher

TCE concentrations.

This study showed that cis-1,2-DCE degradation does not occur to any measurable extent in the

amended anaerobic zone of the study area. Historically, cis-1,2-DCE concentrations in the aquifer

have been less than 50 μ g/L. Therefore, the exact degradation mechanisms for this compound in

this aquifer have not been specifically examined. Continued periodic sampling of wells in the

treatability-study area and in downgradient wells is recommended to understand the fate of this

daughter product. Enhancing the aquifer to create strongly aerobic conditions via sparging may

degrade this daughter product faster in areas where it is now present. This is a possible alternative

for complete treatment of chlorinated compounds at AOC A.

Groundwater samples will be collected from the treatibility study wells and analyzed for

VOC concentrations in July/August 2001, six months after system shutdown, to evaluate further

trends in TCE and daughter product concentrations. Geochemical parameters will continue to be

examined on a biweekly basis for two years to evaluate the pattern of change from anaerobic to

aerobic conditions.

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APPENDIX A WELL CONSTRUCTION LOGS

	A STATE	N	15	5/	7/	-1	=	BOR	ING LOG	of 007G57L	F
)							_				(Page 1 of 2)
		Loca	Millin	ID-SO gton, 1 Buildin	rN. g N-12			Started : 0850 12/1/9 Finished : 0630 12/1/9 Drilling Method : Rotasonic Drilling Company : Boart-Longy Geologist : Bart Dougla	ear	Northing Easting TOC Elevation Total Depth Well Screen	: 392231.80 : 813751.34 : 283.17 : 75 feet : 42 to 72 feet
ł		FI	Jeor #		, 0034		\neg				
	Depth in Feet	in Elev. 283.17 W Rec- FID OF OVERLY (ppm) 5 OF OF OTHER PROPERTY.						DESCRIPTI	l l	007G57LF 283.17 — Cover	
	0		1	100	0			CONCRETE CLAYEY SILT (0.5-6) Dark-brown, moist, and stif	f.		
	10		2	100	0			(6-15) Brown, medium to stiff, and	wet.		
	20-		3	100	0		ML	(15-26) Dark-gray and medium.			High-solids Bent. Grout
7G57LF.BOR	25 -							(26-29) Gray and soft.			
N:WELL LOGS! NSAMIDSOUTH007G57LF.BOR	30-		4	100	0			(29-32) Mottled brown and gray w throughout.	vith a trace of fine	sand	
02-28-2000 N:WELL LOGS!			5	100	0		sc	(32-34) Mottled brown to gray and (34-42) Reddish-brown to orange medium-grained with some clay.		N	
82.28	40	<u> </u>	6	100	0.9		1_			1 4	Bentonite Chip Seal

	ENSAFE							BORING LOG of 007G57LF				
										·	(Page 2 of 2) : 392231.80 : 813751.34 ation : 283.17 th : 75 feet en : 42 to 72 feet Well: 007G57LF Elev:: 283.17	
	NSA MID-SOUTH Millington, TN. Location: Building N-126 Project #: CTO 0094							Started Finished Drilling Method Drilling Company Geologist	: 0850 12/1/99 : 0630 12/1/99 : Rotasonic : Boart-Longyear : Bart Douglas	Total Depti	: 813751.34 ation : 283.17 h : 75 feet	
	Depth in Feet	Surf. Elev. 283.17	SAMPLES	% Rec- overy	FID (ppm)	GRAPHIC LOG	SOIL CLASS	ı	DESCRIPTION	L L		
	40		6	100	0.9		sc					
	1		7	100	1.4	///	SP	SAND, reddish-brow	n to tan and fine-grained			
	45 –		8	100	1.1	0000		SANDY GRAVEL, w medium sand.	rell graded, reddish-brown	n with		
	50		9	100	0.65	0.0.0	GW					
	-		10	100	1	0.00			(Page 2 of 2) : 0850 12/1/99			
	55 - - -		11	100	1.3			GRAVELLY SAND, coarse-grained with	well graded, tan, and me gravel content varying th	dium- to roughout.	(Page 2 of 2) hing : 392231.80 ing : 813751.34 : Elevation : 283.17 il Depth : 75 feet I Screen : 42 to 72 feet Well: 007G57LF Elev.: 283.17	1
	60 –		12	100	0.96			·		·	Sand Pack	
			13	100	1.2		SW		n			
S.	65-		14	100	2.5							
7G57LF.BI	-		15	100	1.1							
SOUTH1007	70-		16	100	0	0.0	GW	SANDY GRAVEL, V	well graded with coarse sa	and.		
N.WELL LOGS! NSAMIDSOUTH:007G57LF.BOR	-		17	100	0		ML	CLAYEY SILT, dark specks throughout.	c-brown to gray with mica	a and lignite		
	75					<u> </u>	<u> </u>	Cockfield contact a	t 72 feet.			
02-25-zbe	80-							1				

	E	Λ	15	5/	7.	F	E	BORI	NG LOG of 00	7G58LF	-
										((Page 1 of 2)
<i>'</i> [Loca	Millin	ID-SO gton, 1 Buildin	rn. g N-12			Started : 0930 11/22/5 Finished : 0910 11/23/5 Drilling Method : Rotasonic Drilling Company : Boart-Longye	99 Easti TOC ear Total	ing Elevation I Depth	: 392157.47 : 8137755.32 : 283.22 : 80 feet
}		Pro	ject #	: CTC	0094	-		Geologist : Bart Douglas	Well	Screen	: 42 to 72 feet
	Depth in Feet	Surf. Elev. 283.22	SAMPLES	% Rec- overy	FID (ppm)	GRAPHIC LOG	SOIL CLASS	DESCRIPTIO	ON	Well: 00 Elev.: 2	07G58LF 83.22 - Cover
	0-							CONCRETE			,
	,		1	100	0		GP	GRAVEL and SAND FILL CLAYEY SILT (2-5) Dark-brown, dry, and stiff.			
	5-1							Wet at 7'.			
	10-		2	100	0			(5-15) Brown, and medium to soft.			
)	15-						ML				
	-						,				— High-solids Bent. Grout
	20-		3	100	0			(15-25) Reddish-brown w/ trace fine medium to stiff.	e sand, moist, and		
								÷ .			
N:NWELL LOGS! NSAMIDSOUTH1007G58LF.BOR	25 <u> </u>		4	100	0		CL	SANDY CLAY, reddish-brown, moi sand.	st and stiff with fine		
OUTHOO!	30-		5	100	2			CLAYEY SAND			
SAMIDS			6	100	1.6			(29-35) Reddish-brown and fine-gra	ained with silt.		
LLOGSI			7	100	2		sc				
TOO N:IMEL	35-		8	100	16.7			(35-39) Changing to tan and yellow	vish-brown.		
25-50	40-	<u> </u>	9	100	27		SM				Bentonite Chip Seal

•

	E	N	15	3/	1	F	E		BORING LO	OG of 007G	558LF	
											(P	age 2 of 2)
		Loca	Millin					Started Finished Drilling Method Drilling Company Geologist	: 0930 11/22/99 : 0910 11/23/99 : Rotasonic : Boart-Longyear : Bart Douglas	Northing Easting TOC Eleva Total Depti Well Scree	h	: 392157.47 : 8137755.32 : 283.22 : 80 feet : 42 to 72 feet
	Depth in Feet	Surf. Elev. 283.22	SAMPLES	% Rec- overy	FID (ppm)	GRAPHIC LOG	SOIL CLASS		DESCRIPTION		Well: 0070 Elev.: 283	
	40-		9	100	27			SILTY SAND, light-be medium-grained with	rown to tan and fine- to a trace of clay.			. •
	- - -		10	100	102		SM		·			
	45-	•	11	100	104			GRAVELLY SAND,w (45-49) Tan to gray a trace of gravel.	ell graded. and fine- to medium-grain	ned with a		
	50		12	100	100			(49-55) Tan and med	lium-grained with gravel	throughout.		
) '	55_		13	100	136		sw			 		
	-		14	100	36 47					·		Sand Pack
	60-		16	100	103			(55-65) Light-brown and gravelly	to tan, medium- to coars	e-grained		0.010 Slotted Screen
	-		17	100	213			"				
SLF.BOR	65		18	100	13	0000		SANDY GRAVEL, w sand.	ell graded and brown wit	h coarse		
UTH1007G5	70-		19	100	39	0.0.	GW					
SAMIDSO	'0-		20	100	35		-	SANDA SILL 1995	addich brown to array			
TL LOGS! A	75-		21	0.	NA				eddish-brown to gray no recovery from 72 to 80) feet		
02-28-2000 N:WELL LOGS! NSAMIDSOUTHOOTGSBLF.BOR	-		22	0	NA.		ML	samples of sandy sill of the sampling rod. estimated at 72 feet.	t were smudged on the be Therefore, the Cockfield terials collapsed and bac	oottom 8 feet I contact was		Backfill
ğ	80-	1	<u> </u>		<u></u>			DOLLOTH S 1881 OF UNITED				

Started 10 10 10 10 10 10 10 1	· }	E	Λ	15	5/	1	F	E	
Coation Building N-126	<i>!</i>		N					<u>-</u>	Finished : 1500 11/29/99 Easting : 813790.26
1 100 2.3									Drilling Company : Boart-Longyear Total Depth : 75 feet
1	-	in Feet	Elev.	SAMPLES	Rec-		GRAPHIC LOG	SOIL CLASS	DESCRIPTION Elev.: 283.17
10 2 35 0 (5-22) Medium to stiff. 3 100 2.2 4 100 0 5 100 0.7 (22-25) Reddish-brown to orange-brown, moist, and medium.		0-		1	100	2.3		GP	CONCRETE GRAVEL and SAND FILL CLAYEY SILT
3 100 2.2 4 100 0 5 100 0.7 (22-25) Reddish-brown to orange-brown, moist, and medium.		5-	·						(2-5) Dark-prown, dry, and very stim.
15	,	10-		2	35	0			(5-22) Medium to stiff.
3 100 2.2 4 100 0 5 100 0.7 (22-25) Reddish-brown to orange-brown, moist, and medium.)	15						ML	
20 4 100 0		-		3	100	2.2			
25 -		20		4	100	0			High-solids Bent. Grou
SANDY CLAY, reddish-brown and stiff with fine sand and silt. CL SANDY CLAY, reddish-brown and stiff with fine sand and silt. CLAYEY SAND, reddish-brown. (28-32) Fine- to medium- grained.		25		5	100	0.7			(22-25) Reddish-brown to orange-brown, moist, and medium.
T 100 4.9 CLAYEY SAND, reddish-brown. (28-32) Fine- to medium- grained. Clayer Sand Cl	7G59LF.BOR	20-		6	100	4.4		CL	silt.
8 100 6.1 SC (32-35) Medium-grained.	AMIDSOUTHIOD	30-		7	100	4.9			
	WELL LOGS! NS	35_		8	100	6.1		SC	

	E	V	15	5/	1	F	E		BORING LO	OG of 0070	659LF	
	سند						-				(1	Page 2 of 2)
<i>j</i>		Loca	Millin	IID-SO igton, Buildin	TN. ig N-1:			Started Finished Drilling Method Drilling Company Geologist	: 0930 11/23/99 : 1500 11/29/99 : Rotasonic : Boart-Longyear : Bart Douglas	Northing Easting TOC Elevi Total Depi Well Scree	th	: 392163.98 : 813790.26 : 283.17 : 75 feet : 42 to 72 feet
			•					<u> </u>				
	Depth in Feet	Surf. Elev. 283.17	SAMPLES	% Rec- overy	FID (ppm)	GRAPHIC LOG	SOIL CLASS		DESCRIPTION		Well: 007 Elev.: 28	1
	40 –		10	100	34	///	sc	(40-42) Reddish-bro	wn to tan.			
	- - -		11	100	44	//,	SP	-	o tan and fine-grained.			-
	45		12	100	50		sw	GRAVELLY SAND, medium-grained.	well graded, light-brown	to tan and		,
	50-	50 - 13 100 69						SANDY GRAVEL, w coarse-grained sand	vell graded with medium- i.	to		
}	55-		14	100	130	0000	GW					
	35-		15 16	100	55 19	0.00						-Sand Pack
	60-		17	100	42		sw	GRAVELLY SAND,	well graded and medium	n-grained.		- 0.010 Slotted Screen
	-		18	100	117	00.00		SANDY GRAVEL V	vell graded with medium	cand		
07G59LF.BOR	65-		19	100	33	0.00	, CW	OAND TOTALL, V	ren gradea wan mediam	Janu.		
N.WELL LOGS! NSAMIDSOUTH/007G59LF.BOR	70-		20	100	38	0000			-			
ELL LOGS! N	75-		21	100	3.4		ML	CLAYEY SILT, dark lignite throughout. Cockfield contact at	e-gray to brown, micacius	s and soft with		
W:W				· · · · · · · · · · · · · · · · · · ·						/		

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	E	N	15	5/	7/	F	E		BORING LO	G of 007G60	LF
	سعمد،		_	-							(Page 1 of 2)
<i>/</i> [Loca	Millin	ID-SO gton, Buildin	TN. g N-12			Started Finished Drilling Method Drilling Company Geologist	: 0730 11/30/99 : 2200 11/30/99 : Rotasonic : Boart-Longyear : Bart Douglas	Northing Easting TOC Elevation Total Depth Well Screen	: 392114.00 : 813797.36 : 283.26 : 85 feet : 45 to 75 feet
			,,001 11		0001		Ī			1	
	Depth in Feet	Surf. Elev. 283.26	SAMPLES	% Rec- overy	FID (ppm)	GRAPHIC LOG	SOIL CLASS		DESCRIPTION		007G60LF : 283.26 — Cover
	0-							CONCRETE			
	-		1	40	0			CLAYEY SILT (1-6.5) Dark-brown,	moist, and medium.		
	5-		2	100	3.2			·			
	10		3	100	0.8			(6.5-19) Brown, soft feet.	and wet. Medium stiffnes	ss below 16	
)	15		4	100	3.4		ML		•		
			5	100	2.2				-		
	20-			100	6.8			(19-23) Reddish-bro	own and medium.		High-solids Bent. Grout
	25		7	100	5			(23-25) With fine sa		ın and	
œ			8	100	9.9		CL	medium with fine sa	dish-brown to orange-brow and	in allu	
7.G80LF.BO	30-		9	100	3.7				own to orange-brown, fine	- to	
IDSOUTHOO	35-		10	100	9.3		sc S				
N:WELL LOGS! NSAMIDSOUTH1007G60LF.BOR	-		11	100	12						
			12	100	24			with some clay.	own to tan and fine- to med		Bentonite Chip Seal
02-20-200	45-		13	100	38		SF		to tan and fine- to medium	n-grained.	Sand Pack

	E	Λ	15	5/	1	F	É		BORING LO	OG of 007G	
/		N		IID-SC ngton,			-	Started Finished	: 0730 11/30/99 : 2200 11/30/99	Northing Easting	(Page 2 of 2) : 392114.00 : 813797.36
				Buildir #: CTC	_			Drilling Method Drilling Company Geologist	: Rotasonic : Boart-Longyear : Bart Douglas	TOC Eleva Total Depti Well Scree	1 : 85 feet
ł			JCCL F	. 010	000-	,		1 000,000			
	Depth in Feet	Surf. Elev. 283.26	SAMPLES	% Rec- overy	FID (ppm)	GRAPHIC LOG	SOIL CLASS		DESCRIPTION	•	Well: 007G60LF Elev.: 283.26
	45_		14	100	25		SP	GRAVELLY SAND),well graded.	e of gravel.	
	50		15	100	57			(10 00), 1211 2112 111			
	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		16	100	71			(50-65) Tan and m	nedium- to coarse-grained	with gravel	
	55_		17	100	37		sw	content varying the	roughout.		: : : : : :
	60-		18	100	43						0.010 Slotted Screen
	65-		19	100	55						Sand Pack
	05-		20	100	NA			(65-69) Brown to	tan.		
	70-		21	100	NA	00,0	1	SANDY GRAVEL	, well graded with coarse s	and.	
-			22	100	NA	000	GW				
GEOLF.BOR	75~		23	100	NA		٠.	lignite specks thro	y and clayey with fine sand ughout. One reddish-oran	d, mica, and age iron	
02-28-2000 N:WELL LOGS! NSAMIDSOUTH1007G80LF.BOR	80-		24	0	NA		ML	concreation at 74. Cockfield contact	•		
N:WELL LOGS! N	85-										Bentonite Chip Seal
02-28-2000	90-							•			

E	Λ	15	5/	1	F	E	BORING LOG	of 007	'G61LF	•
							·		((Page 1 of 2)
	Loca	Millir	ngton, Buildir	TN. ng N-1			Started : 1000 12/2/99 Finished : 1800 12/2/99 Drilling Method : Rotasonic Drilling Company : Boart-Longyear Geologist : Bart Douglas	Easting TOC EI Total Do	evation epth	: 392122.48 : 813813.14 : 283.04 : 75 feet : 45 to 75 feet
Depth in Feet	Surf. Elev. 283.04	SAMPLES	% Rec- overy	FID (ppm)	GRAPHIC LOG	SOIL CLASS	DESCRIPTION			
0-11111		1	100	0			CONCRETE CLAYEY SILT (1-4) Dark-brown.			
10		2	100	0		ML	(4-16) Brown.			
20-		3	100	0			(16-25) Reddish-brown.			- High-solids Bent. Grout
25		4	100	0		CL	CLAYEY SAND, reddish-brown to orange-brown and			·
35_		5	100	0 5.6		sc	SAND, yellowish-brown to tan and fine- to			•
	Depth in Feet	Depth in Elev. Feet 283.04	NSA M Milling Incomplete Incomplete	NSA MID-SC Millington, NSA MID-SC Millingt	NSA MID-SOUTH Millington, TN.	NSA MID-SOUTH Millington, TN.	NSA MID-SOUTH Millington, TN.	NSA MID-SOUTH Millington, TN. Started :1000 12/2/99 Filling Method Rotasonic Drilling Method Rotasonic Drilling Company :Boart-Longyear Geologist :Bart Douglas Sandard Filling Method Rotasonic Drilling Company :Boart-Longyear Geologist :Bart Douglas Sandard Filling Company :Boart-Longyear Geologist :Bart Douglas Sandard :1000 12/2/99 Filling Company :Boart-Longyear Geologist :Bart Douglas :Bart	NSA MID-SOUTH Millington, TN. Started :1000 12/2/99 Easting Drilling Method Rotazonic TOC E Easting Drilling Method Rotazonic TOC E Color Easting Drilling Method Rotazonic TOC E Color Easting Drilling Method Rotazonic TOC E Color Easting Drilling Company Boart-Longyear Total D Geologist Earl Douglas Well Sc Easting Color Easting Drilling Company Boart-Longyear Total D Color Easting	NSA MID-SOUTH Millington, TN.

	E	N	15	3/	1	F	E		BORING LO	OG of 00		
4				IID-SO			· 	Started	: 1000 12/2/99	Northi	ng .	(Page 2 of 2)
		Loca	ation:	gton, Buildin	g N-1			Finished Drilling Method Drilling Company Goologist	: 1800 12/2/99 : Rotasonic : Boart-Longyear : Bart Douglas	Eastin TOC E Total I Well S	Elevation Depth	: 813813.14 : 283.04 : 75 feet : 45 to 75 feet
-		Pro	oject #	E CTC	0094			Geologist	: Bart Douglas	vven	Ceen	. 45 to 75 leet
	Depth in- Feet	Surf. Elev. 283.04	SAMPLES	% Rec- overy	FID (ppm)	GRAPHIC LOG	SOIL CLASS		DESCRIPTION		Well: 00 Elev.: 2	97G61LF 83.04
	40_		6	100	5.6							- High-solids Bent. Grout
			7	100	182		SP					– Bentonite Chip Seal
	45-		8	100	57			GRAVELLY SAND,	well graded and mediur	n-grained.		
	50-		9	100	13		sw					
	50-		10	100	93	0.00		SANDY GRAVEL, v	well graded with medium	to coarse		
	- - - 55-		11	100	285	0.00		Saliu.		·		
	55 -		12	100	85	0.00						
			13	100	78	0.0.		•				0.010 Slotted Screen
	60-		14	100	37	0000	1		·			— Sand Pack
	65 -		15	100	78	0000	GW					
G61LF.BOR			16	100	27	0.0						
COUTHIOO?	70 -					0.00						
3SI NSAMIDS			17	100	18	0.0	,					
בר ומ	75-]					ML	CLAYEY SILT, gra	y with mica throughout.	•	1	}
W.K		1						Cockfield contact a	at 74 feet.			
02-28-200- N:WELL LOGS! NSAMIDSOUTH1007G61LF.BOR	80-	1						L			_1	

	Λ						LOG OF BORII			(Page 1 of 2)
		Millin	IID-SC igton, Buildir		26		Started : 1100 8/3/00 Finished : 1500 8/3/00 Drilling Method : Rotasonic Drilling Company : Alliance Environmental	Northing Easting TOC Ele Total De	vation	: : 75 feet
				0 0094		···	Geologist : Bart Douglas	Well Scre		: 45 to 75 feet
Depth in Feet	Surf. Elev.	Samples	% Rec- overy	FID (ppm)	GRAPHIC	nscs	DESCRIPTION		Well: 00 Elev.:	7G62LF - Cover
0	- 0	1	100	0			CONCRETE CLAYEY SILT			
5-	5		100	,			(1-6.5) Dark-brown, moist, and medium stiff.			
-		2	100	0						,
10	- - 10	3	100	0			(6.5-19) Brown, soft, and wet. Medium stiffness be 16 feet.	elow		
15	15		100			ML				
1										
20	- -2 0	4	100	0			(19-23) Reddish-brown and medium stiffness.			-High-solids Bentonite Grout -2" PVC Riser
25	25						(23-25) With fine-grained sand throughout. SANDY CLAY, brown to orange-brown and stiff with	th		
- - - - - - - - - -						CL	very-fine- to fine-grained sand.			
30	30	5	100	0						
35	35						CLAYEY SAND, (33-38) Orange-brown and fine-grained.	<u>. : · </u>		·
- - - - - - -		6	100	NA		sc	(38-42) Yellowish-brown to tan and very-fine- to fine-grained.			

	E	7	15	5/	7	F	E		LOG OF BORI	NG 007G	62LF	:
				77			-				(F	age 2 of 2)
		Loca	Millin	IID-SO igton, ⁻ Buildin #: CTO	ΓΝ. g N-1			Started Finished Drilling Method Drilling Company Geologist	: 1100 8/3/00 : 1500 8/3/00 : Rotasonic : Alliance Environmental : Bart Douglas	Northing Easting TOC Elevatio Total Depth Well Screen	'n	: : : 75 feet : 45 to 75 feet
	Depth in Feet	Surf. Elev.	Samples	% Rec- overy	FID (ppm)	GRAPHIC	nscs		DESCRIPTION		ell: 007 ev.:	G62LF
	40-	-40		T		///	sc					
	-		6	100	NA	///		SAND, Poorly Grade	ed, tan, and fine- to medium-	grained.	排	Bentonite Chip Seal
	45_	-45					SP	At 42 is a 1/2-inch th	ick pinkish-gray clay seam.			2" PVC Riser
	50-	-50	7	100	NA.		sw	SAND, Well Graded coarse-grained with	, tan to light-brown and medi gravel.	um- to		
)	-					.0.0		GRAVEL, Well Grac coarse-grained sand	led, and gray with medium- to	0		
	55 <u> </u>	-55					GW					Sand Pack
	60-	-60	8	100	NA		SW	SAND, Well Graded	l, tan, gravely, and coarse-gr	ained.		0.010 Slotted Screen
32LF.BOR	65 -	-65	-					GRAVEL, Well Grad	ded, with coarse-grained san	d .		
N.WELL LOGSUNSAMIDSOUTH/007G62LF.BOR	70-	-70	9	100	NA		GW					
N:WELL LOGSW	75 -	-75						CLAYEY SILT,				
07-02-2001	İ						ML	At 75 feet hit 1 inch and orange-brown of Cockfield.	of mottled light-gray, pinkish clayey silt, with mica flakes. I	-mauve, Fop of		
0.20	80-	1			<u> </u>		Щ			<u> </u>		

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	E	Λ	15	3/	4	F	E		LOG OF BOF	RING 007	G63L	F
						=_						(Page 1 of 2)
		Loca	Millin					Started Finished Drilling Method Drilling Company Geologist	: 1705 8/3/00 : 1830 8/3/00 : Rotasonic : Alliance Environmental : Bart Douglas	Northing Easting TOC Elev Total Dep Well Scre	th	: : : 75 feet : 45 to 75 feet
	-											*
	Depth in Feet	Surf. Elev.	Samples	% Rec- overy	FID (ppm)	GRAPHIC	nscs		DESCRIPTION		Well: 00 Elev.:	97G63LF - Cover
	0-	- 0		_			I	CONCRETE			TIT	ו
			1	100				CLAYEY SILT (1-7) Brown and me	edium to stiff.			
	5	5										
	10	-10	2	100				(7-10) Dark-brown (10-18) Brown and	and medium to stiff.			
)	15_	-15					ML					
	20-	-20	3	100				(18-25) Brown to o	range-brown and medium.	1		— High-solids Bentonite Grout — 2" PVC Riser
BOR	25	-25							y-fine-grained sand and med			
DSOUTH007G83LF.	30-	-30	4	100			CL	·	ttled red-brown and orange- grained sand. eddish-brown, fine-grained, a			
N:WELL LOGSINSAMIDSOUTH007G83LF.BOR	35-	-35					sc		ng more yellowish to mustaro			
07-02-2001			5	100								— Bentonite Chip Seal

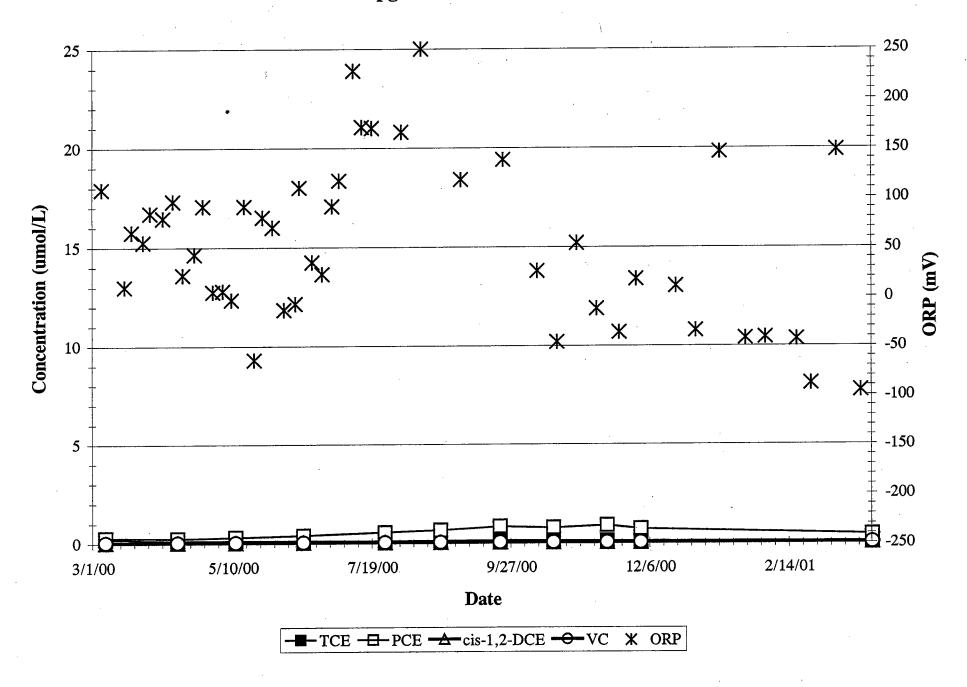
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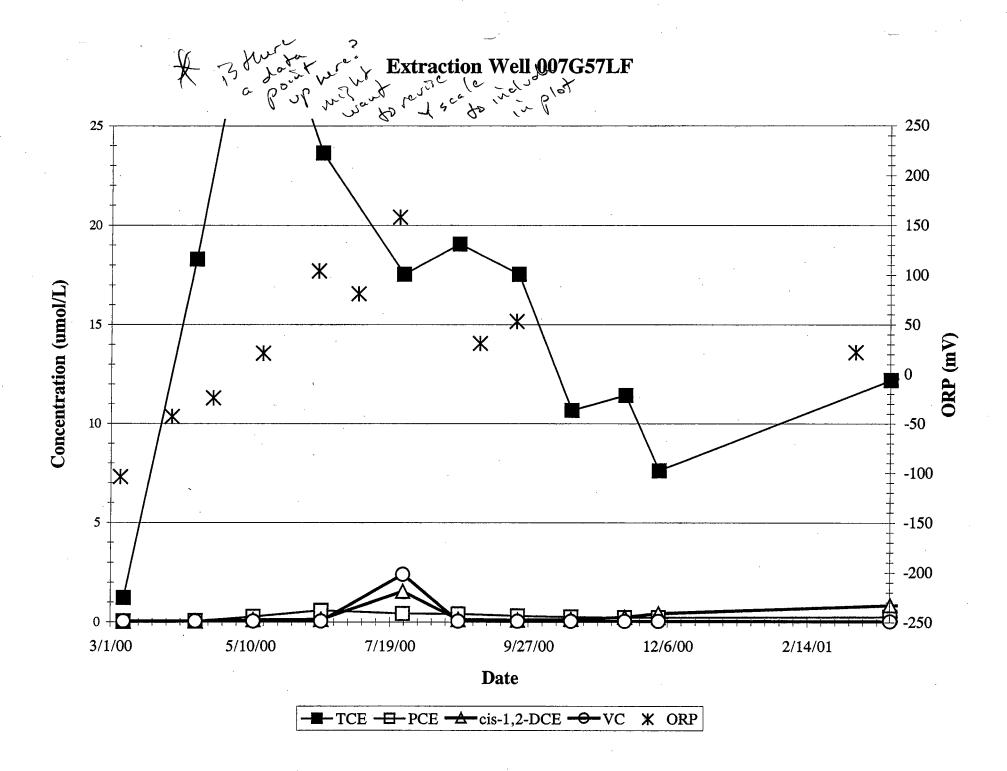
•

	E	7	15	5/	4	F	E		LOG OF BORI	NG 007	G63L	F		
	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,											(Page 2 of 2)		
		Loca	Millin	IID-SC igton, ' Buildir #: CT(TN. ig N-12			Started Finished Drilling Method Drilling Company Geologist	: 1705 8/3/00 : 1830 8/3/00 : Rotasonic : Alliance Environmental : Bart Douglas	Northing Easting TOC Eleve Total Depl	th	: : : 75 feet : 45 to 75 feet		
		PI	oject	#: 010	J 0092	•		Coologist	. Dan Douglad			. 70 10 10 100		
	Depth in Feet	Surf. Elev.	Samples	% Rec- overy	FID (ppm)	GRAPHIC	nscs	D	ESCRIPTION	4	Well: 00 Elev.:	07G63LF		
	40-	-40							hick, light-gray silty clay sea		MA	Bentonite Chip Seal		
			5	100			SIVI		and very-fine- to fine-graine , tan to light-gray and very-t			– 2" PVC Riser		
	45-	-45					sw	SAND, Well Graded, t with some gravel.	SAND, Well Graded, tan, and medium- to coarse-grained vith some gravel.					
	50	-50	6	100		. 0 . 0 .		GRAVEL, Well Grade	d, with medium-grained san hick light-gray clay seam.	d.				
	 - - -					· · · · · ·		SAND, Well Graded, gravel.	medium- to coarse-grained o					
:	55_ 	-55	7	100			GW	sand.	5, W an moulant to course ,	gramou		— Sand Pack — 0.010 Slotted Screen		
	- -	-				0 0 0	SW GW SW	gravel.	an, and medium-grained w	/				
ĸ	65-	-65				0 0 0 0 0		SAND, Well Graded, gravel. GRAVEL, Well Grade	an, and medium-grained w	rith				
N:WELL LOGSINSAMIDSOUTH1007G83LF.BOR	`70-	-70	8	100		.0.9.	GW	sand.						
LL LOGSYNSAM	75-	-75				0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	ML CL	interbedded thin laver	LTY CLAY, light-gray to lights of silty clay and clayey sil	nt-tan t with				
07-02-2001 N:WE	•							mica flakes. Top of C	осклета.		,			

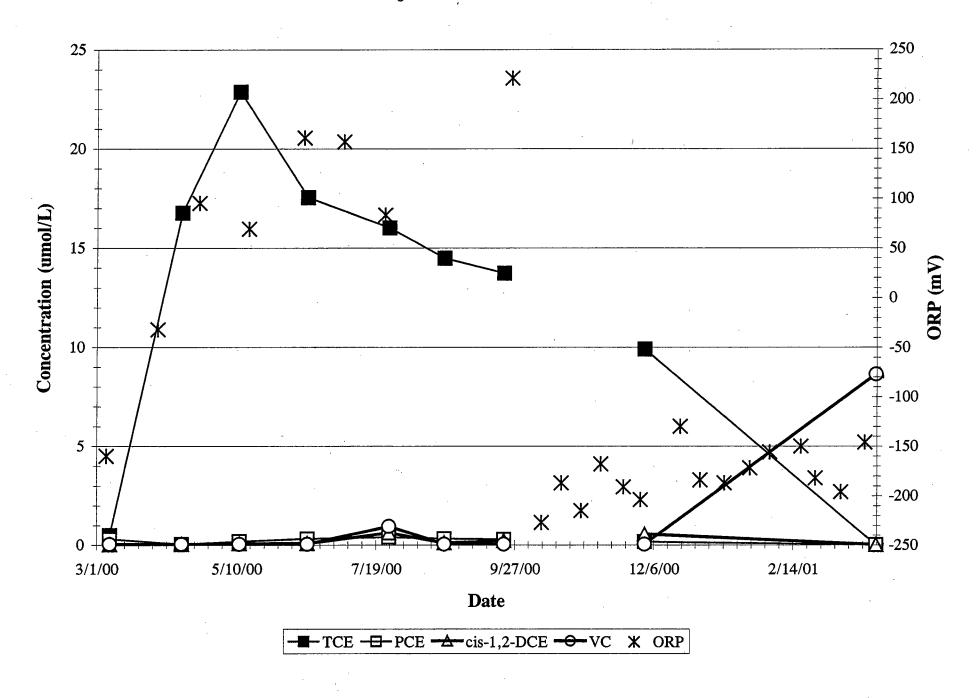
APPENDIX B VOC DATA GRAPHS

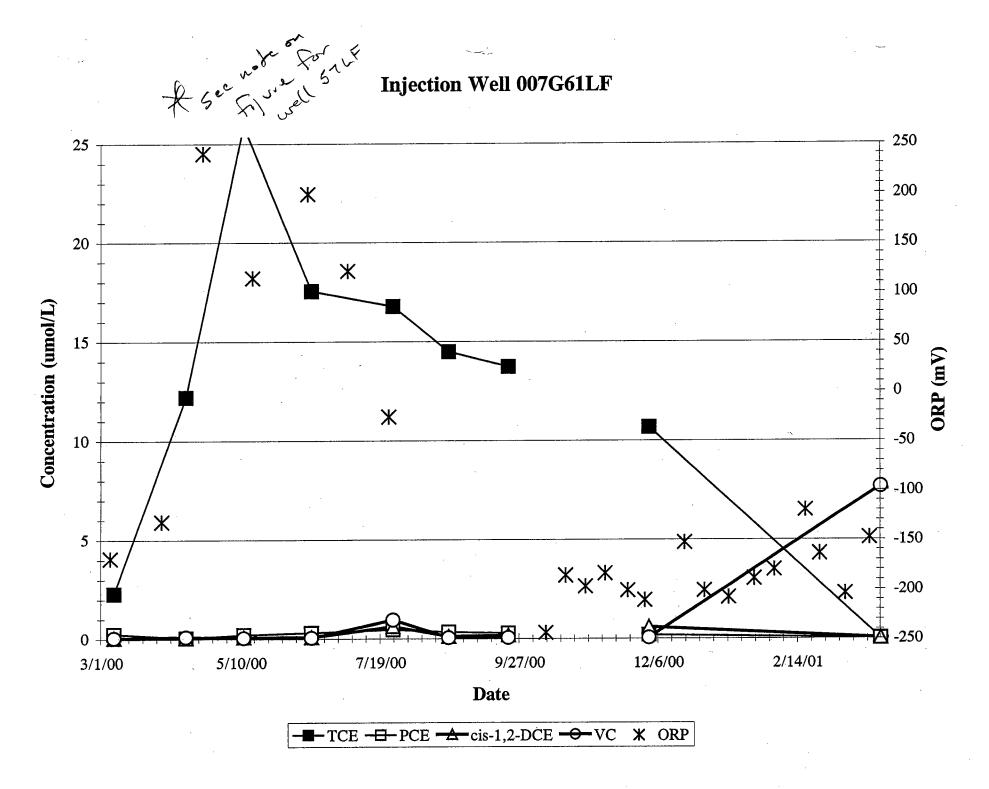
Upgradient Well 007G03LF



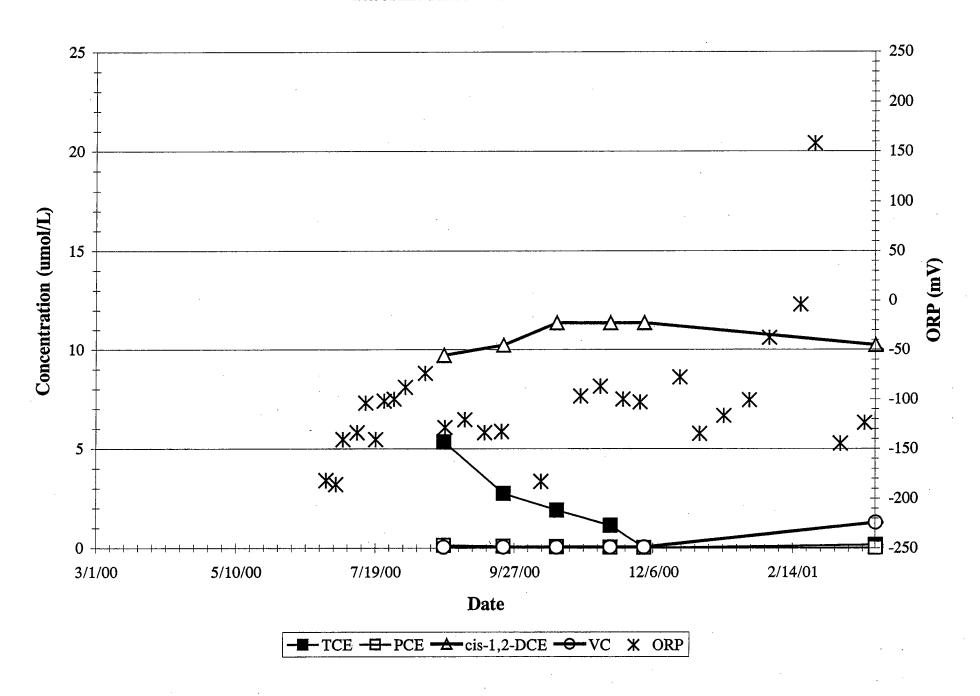


Injection Well 007G60LF

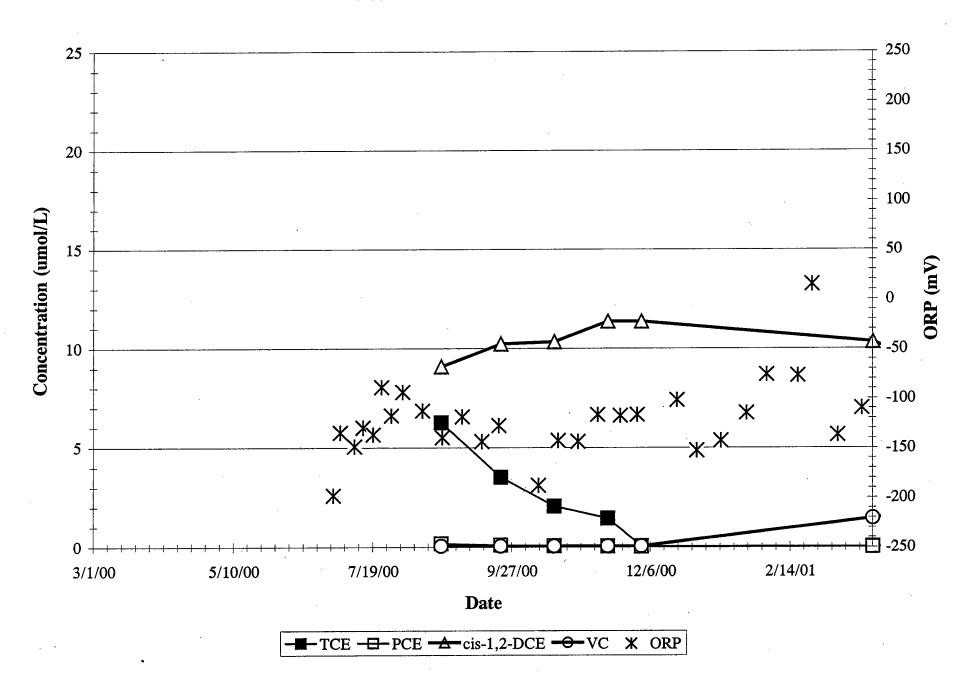




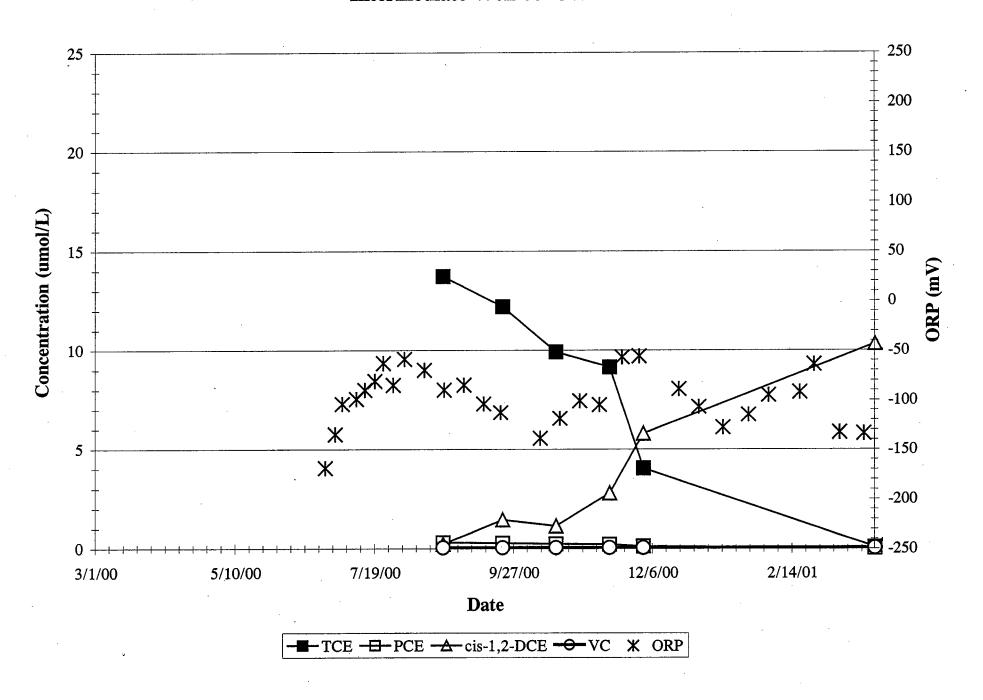
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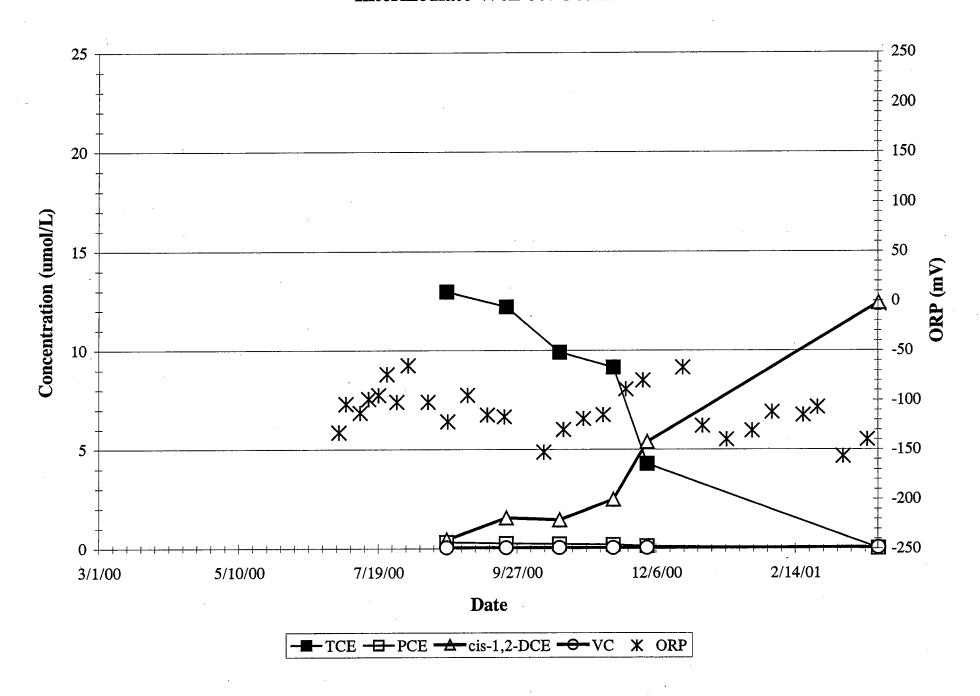
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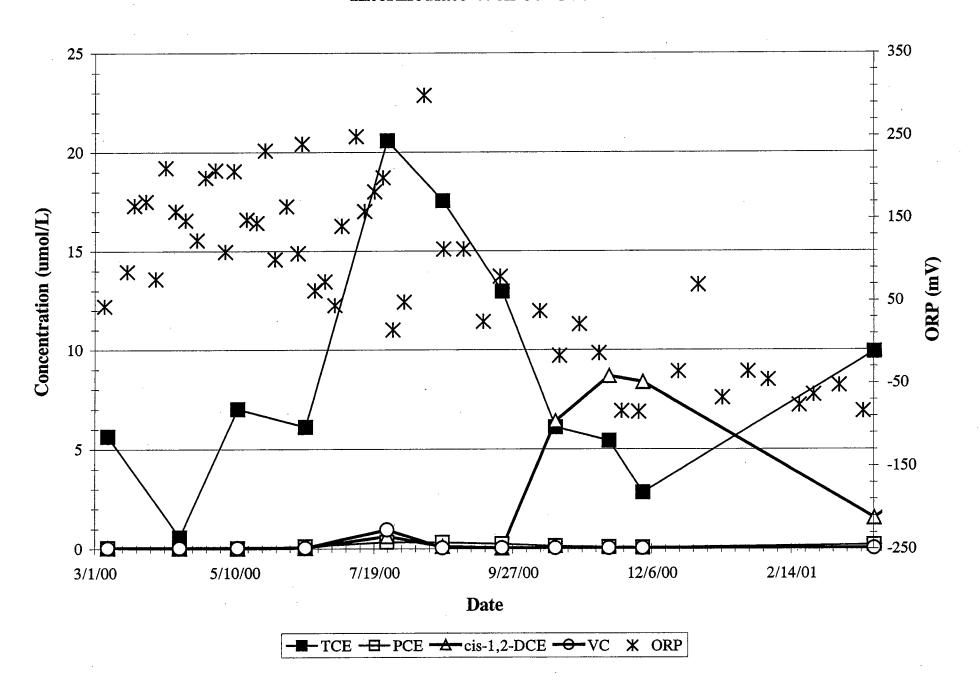
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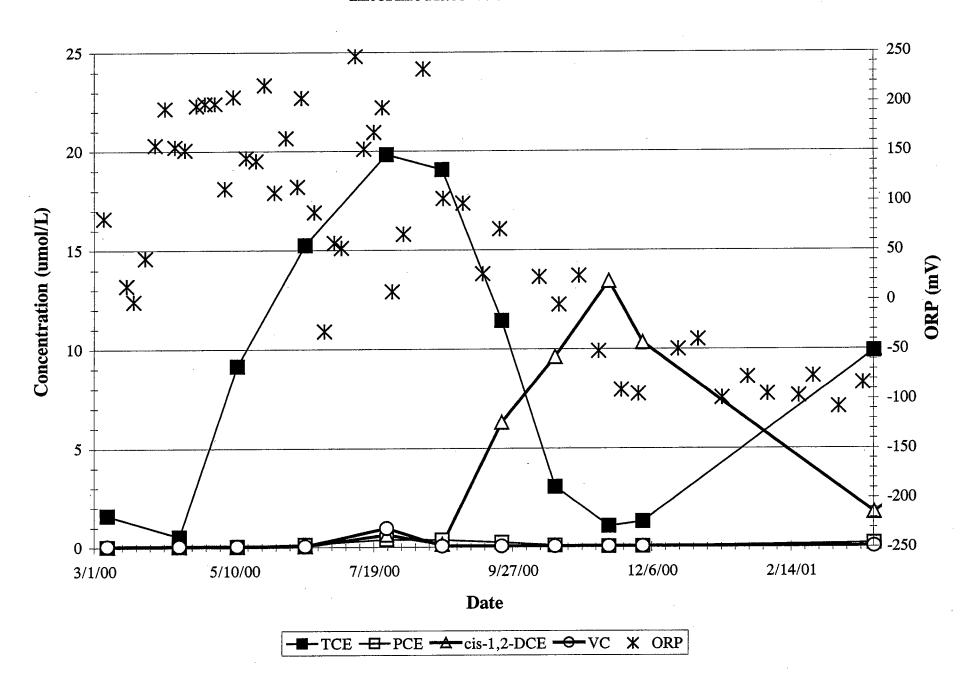
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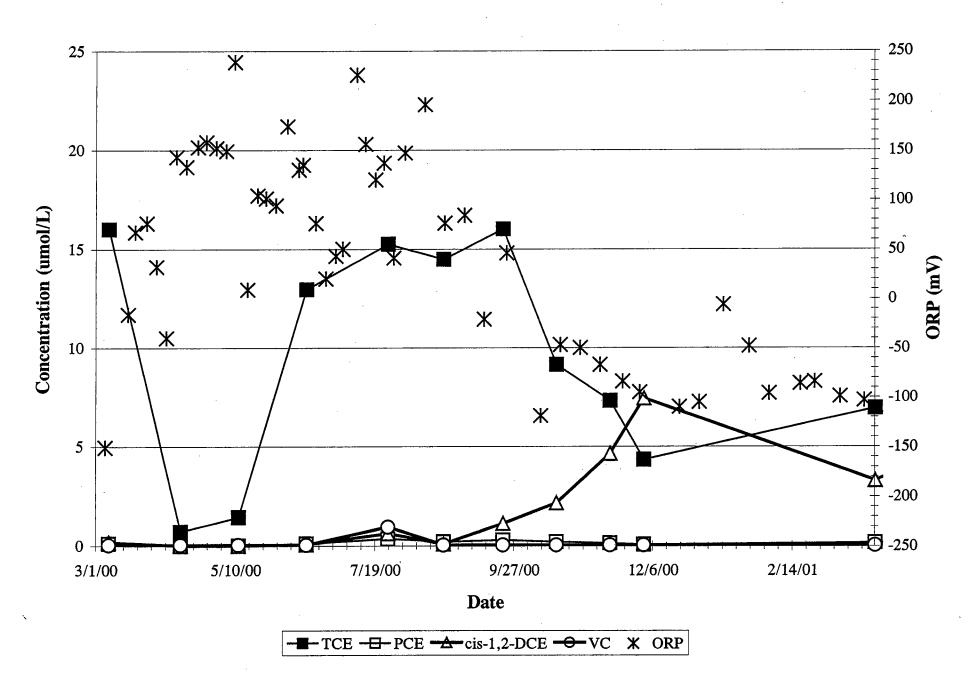
Intermediate Well 007G58LFA



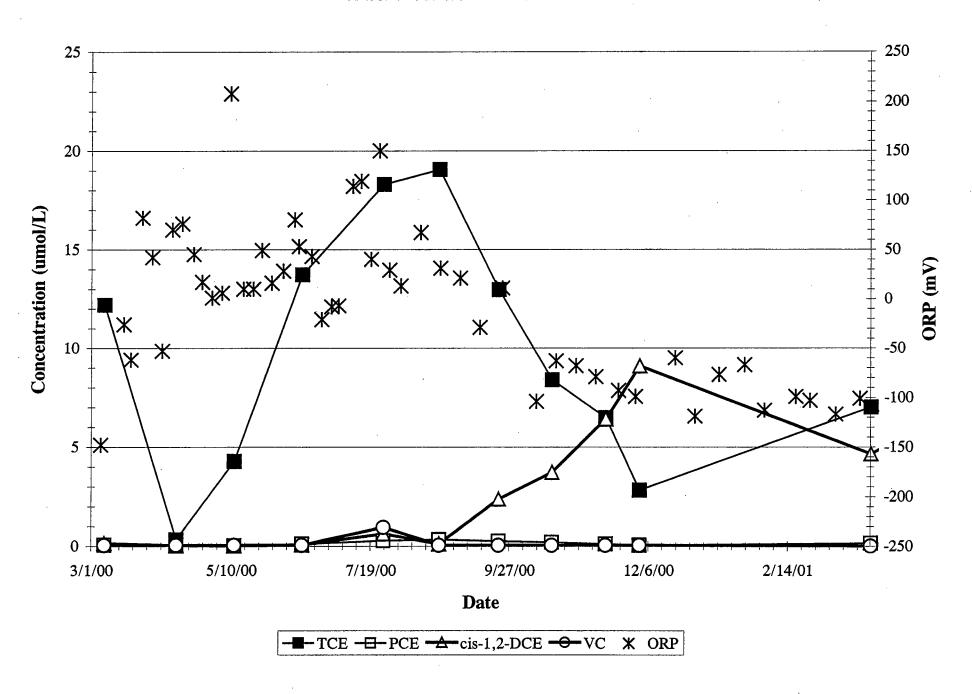
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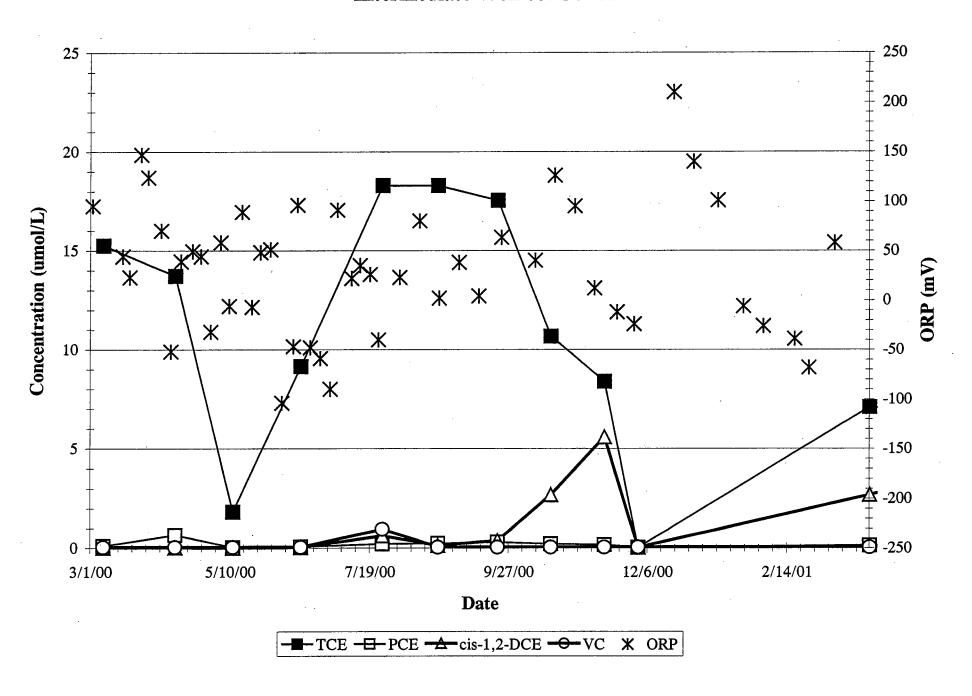
Intermediate Well 007G59LFA



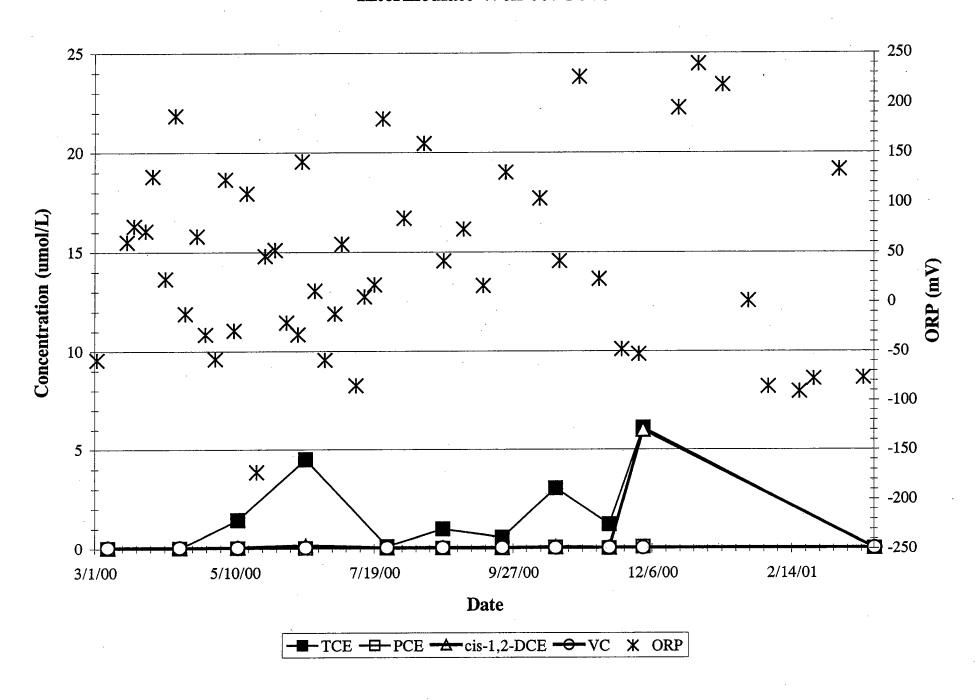
Intermediate Well 007G59LFB



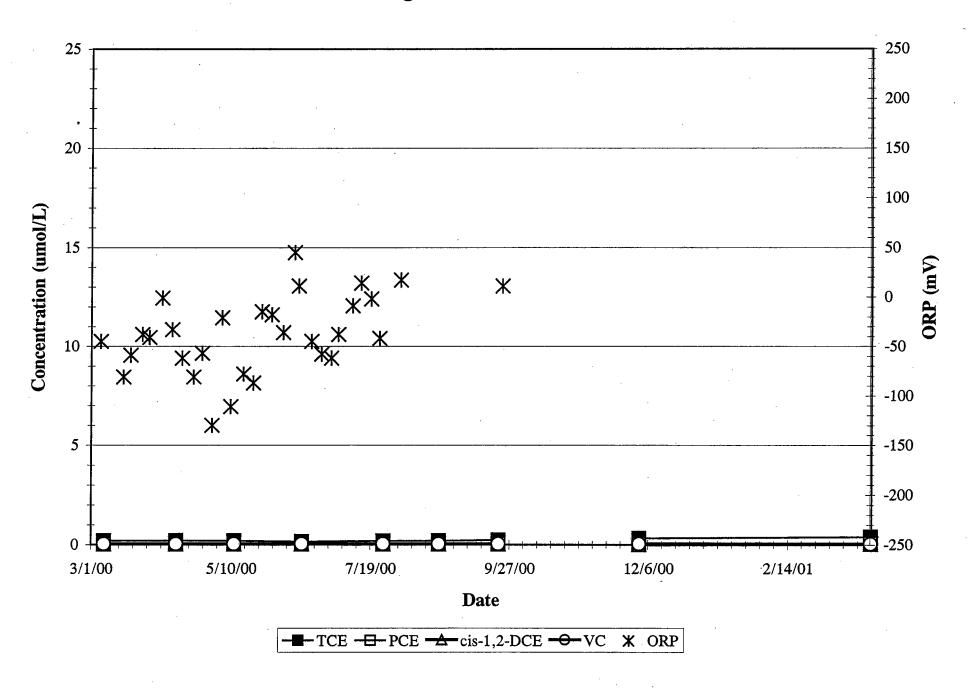
Intermediate Well 007G04LF



Intermediate Well 007G04UF

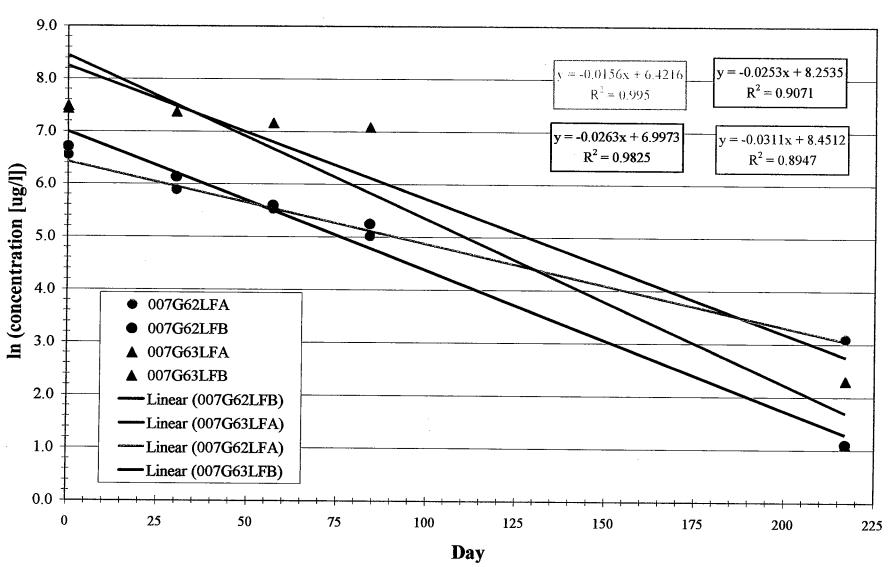


Downgradient Well 007G21LF

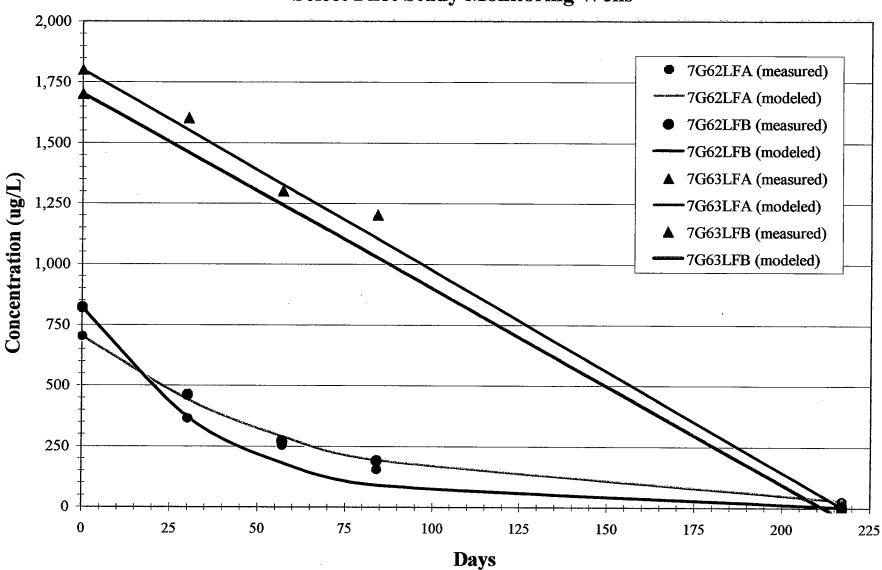


APPENDIX C DEGRADATION RATE GRAPHS

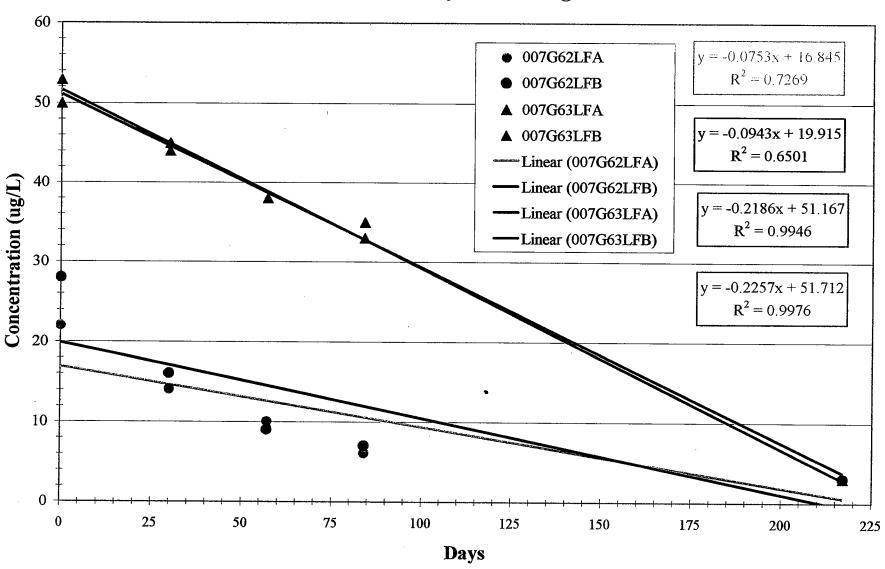
First-Order TCE Degradation Rates for Select Pilot Study Monitoring Wells



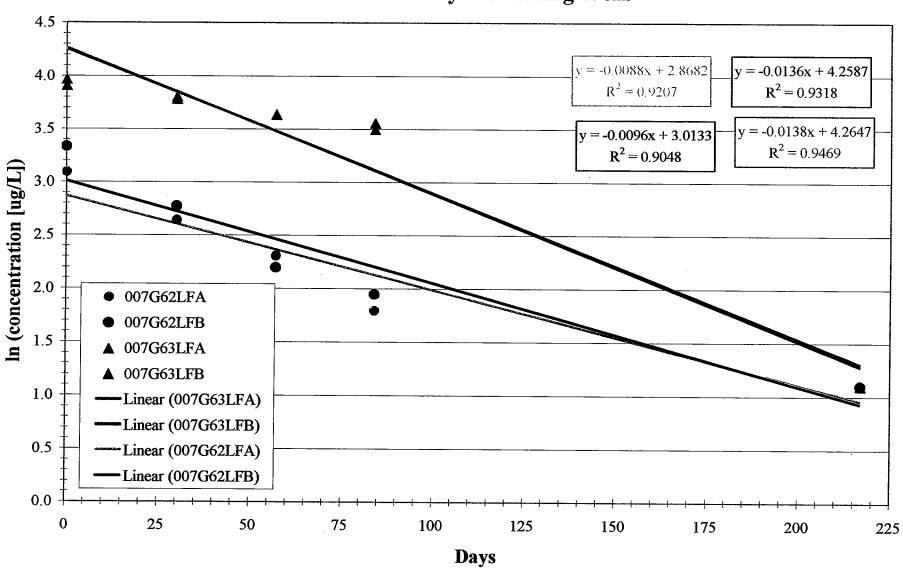
Measured and Modeled TCE Concentrations for Select Pilot Study Monitoring Wells



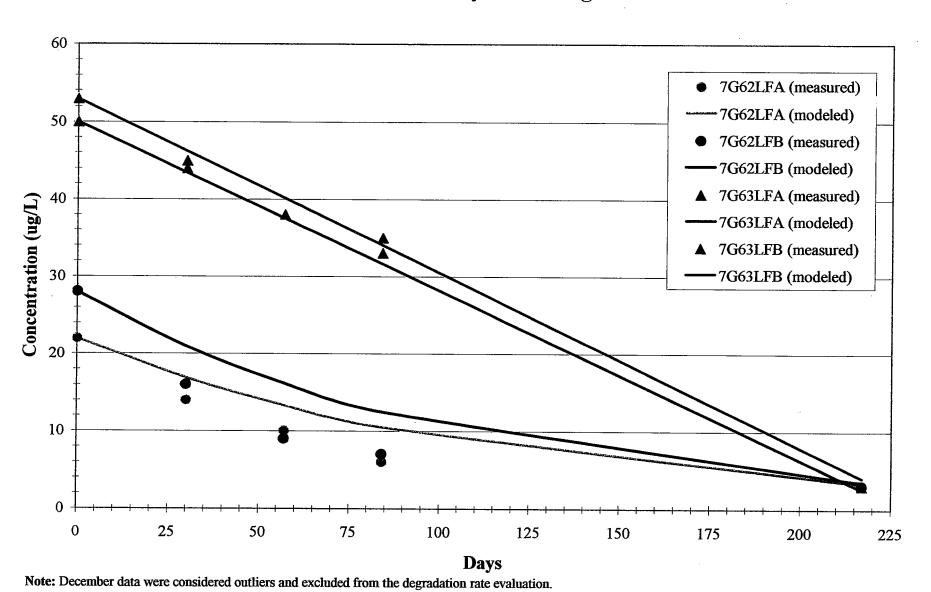
Zero-Order PCE Degradation Rates for Select Pilot Study Monitoring Wells



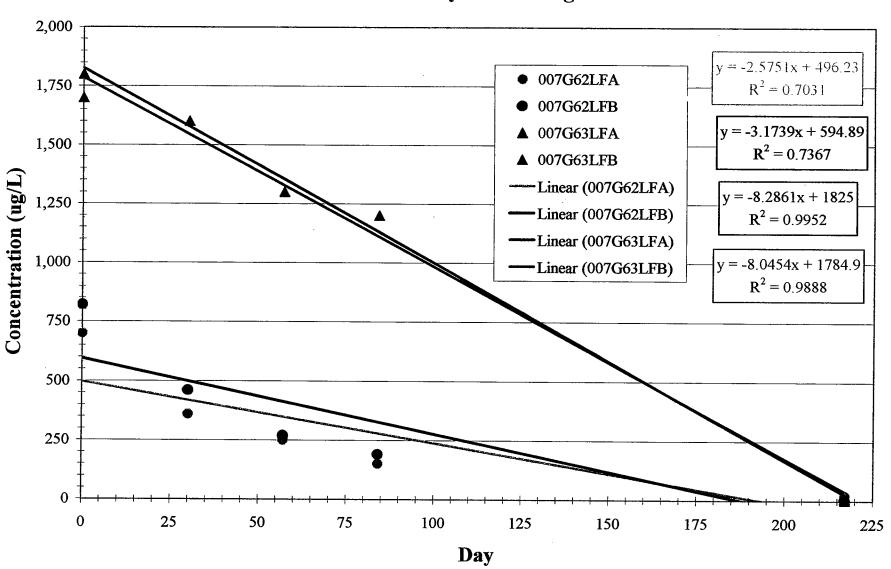
First-Order PCE Degradation Rates for Select Pilot Study Monitoring Wells



Measured and Modeled PCE Concentrations for Select Pilot Study Monitoring Wells



Zero-Order TCE Degradation Rates for Select Pilot Study Monitoring Wells



APPENDIX D GEOCHEMICAL DATA

Field Data

ORP (mg/L) (field)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFR	58LFA	581.FR	591.FA	591.FR	4LF	4UF	21LF
3/2/00								00.537.25	- COZZIII	- COLII D	U) DI II	37212	95	-59	2111
3/7/00	108	-104	-160	-169	,			*	43	82	-151	-148	75	-37	-45
3/23/00	10			-02					85	14	-16	-26	44	60	-81
3/28/00	65								165	-2	67	-62	23	76	-59
4/5/00	55								170	42	76	82	147	71	-38
4/10/00	84								170	725	70	02	124	126	-38 -41
4/12/00		-43	-32	-132					76	156	32	42	124	120	-41
4/19/00	79		32	132					211	193	-40	-53	70	23	-1
4/26/00	96								158	154	143	70	-52	23 187	-33
5/3/00	22								147	151	133	76	39	-12	-62
5/11/00	43	-24	95	240					123	196	153	45	49	-12 66	-02 -81
5/17/00	91	27	75	240					199	198					
5/24/00	5				:				208	198	158	17	44	-33	-57 120
5/31/00	6								109		152	1	-32	-58 122	-130
6/6/00	3								207	112 205	149	.6	58	123	-21
6/15/00	91	21	69	114					1		239	208	6	-29	-111
6/22/00	-64	21	09	114	[148	143	9	10	89	109	-78 -77
6/28/00	80								144	140	104	10	-7 40	-173	-87
7/5/00	70								232	217	101	49	48	46	-15
7/13/00					ŀ				100	108	94	16	51	52	-18
	-13				ŀ				164	163	174	28	-104	-21	-36
7/21/00	-7	104	1.61	100					107	114	130	80	-47	-33	45
7/24/00	110	104	161	199					240	204	135	53	96	141	11
8/2/00	35				100		4.00		62	88	76	43	-48	11	-45
8/9/00	23				-182	400	-169	400	73	-32	20	-21	-59	-59	-58
8/16/00	91	0.4		404	-186	-198	-135	-133	44	57	43	-8	-90	-12	-62
8/21/00	117	81	157	121	-141	-135	-104	-104	140	52	50	-7	91	58	-38
8/31/00	228				-134	-149	-99	-113	249	246	226	114	22	-85	-9
9/6/00	171				-104	-130	-90	-99	158	152	156	119	35	5	14
9/13/00	170				-141	-137	-81	-95	182	169	120	40	. 26	17	-2
9/19/00	277	158	83	-26	-102	-89	-63	-74	199	194	137	150	-40	184	-42
9/26/00					-100	-118	-85	-102	14	8	41	29			
10/4/00	166				-88	-94	-59	-65	48	66	147	13	23	84	17
10/18/00	250	271			-74	-113	-70	-102	299	233	196	67	80	159	
11/1/00					-129	-140	-90	-122	112	102	76	31	2	41	
11/15/00	118	31			-121	-119	-85	-95	112	97	84	21	38	73	
11/29/00					-134	-144	-104	-115	24	26	-21	-29	4	16	
12/11/00		53			-133	-128	-113	-117	79	71					
12/15/00	138	•									46	11	63	130	11
12/18/00			221	262								İ			
1/8/01	26		-227	-244	-183	-188	-139	-153	37	23	-119	-104	40	104	
1/22/01	-46		-187	-186		-143	-119	-130	-17	-5	-47	-63	126	41	
2/5/01	54		-215	-197	-97	-144	-101	-119	21	24	-50	-68	95	226	
2/19/01	-12		-168	-184	-87	-117	-105	-115	-14	-52	-67	-79	12	23	
3/7/01	-36		-191	-201	-100	-118	-57	-89	-84	-91	-84	-93	-12	-48	
3/19/01	18		-204	-211	-103	-117	-56	-80	-85	-95	-95	-99	-24	-53	
					-				•			1		_	

DO (mg/L) (field)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
3/2/00						·							1.50	1.60	
3/7/00												1			
3/23/00	1.90								3.00	3.30	1.30	1.90	1.80	1.00	0.80
3/28/00	2.20			·					3.50	3.40	1.20	1.30	2.00	0.80	0.70
4/5/00	0.50								2.10	2.30	1.70	0.70	1.80		0.50
4/10/00	2.80						-						2.80	0.80	
4/12/00	1.30	1.00	1.20	1.00					3.50	2.20		0.60	2.20		0.70
4/19/00									1.70	2.10	0.80				0.60
4/26/00										1.07		ļ	0.20		•
5/3/00	0.80								1.80	•	0.80			0.80	0.40
5/11/00	0.65	0.83	0.99	0.40]				1.37	0.30	0.64	0.43	0.90	0.69	0.15
5/17/00	1.03				ļ				0.94	0.30	0.31	0.28	0.81	0.10	0.11
5/24/00	2.13				1				1.01	0.30	0.38	0.43	1.65	0.46	0.18
5/31/00	1.50				ł				1.20	0.23	0.43	0.35	1.98	0.15	1.01
6/6/00	1.89								0.72	0.66	0.40	0.12	0.30	0.20	0.12
6/15/00	1.94	0.49	1.86						0.65	0.23			1.38	0.40	1.15
6/22/00	1.46				ľ				0.46	0.09	0.25	0.13	0.45	0.06	0.12
6/28/00	2.02								0.87	0.19	0.47		0.17	0.12	0.14
7/5/00					1	•]						
7/13/00	0.15				Ì				0.42	0.25	0.48	0.12	0.32	0.17	0.13
7/21/00	2.06								0.51	0.19	0.16				
7/24/00	2.25	1.15	2.90	1.72					1.04	0.89	0.75	0.73	1.22	0.60	0.78
8/2/00	1.22								0.51	0.30	0.30	0.17	0.22	0.19	0.21
8/9/00	0.37				0.06		0.12		0.34	0.09	0.19	0.08	0.18	0.15	0.19
8/16/00	0.12				0.06	0.04	0.05	0.06	0.38	0.12	0.23	0.22	0.15	0.16	0.35
8/21/00	2.80	1.83	3.59		ŀ	0.52			1.86	0.61	0.70				1.42
8/31/00	1.85				0.10	0.07	0.13	0.12	0.45	0.16	0.21	0.17	0.15	0.07	0.08
9/6/00	1.33				0.09	0.06	0.08	0.13	0.28	0.11	0.22	0.12	0.14	0.10	0.11
9/13/00	1.75				0.12	0.15	0.15	0.19	0.92	0.25	0.63	0.50	0.19	0.27	0.18
9/19/00	3.01	1.95							1		0.18		2.11		
9/26/00	1.47				0.15	0.12	0.12	0.11	0.34	0.11	0.21	0.16	0.13	0.10	0.14
10/4/00	1.44				0.16	0.11	0.13	0.14	0.32	0.19	0.24	0.18	0.24	0.19	0.21
10/18/00	1.68	0.63			0.54	0.20		0.37	0.85	0.54	0.79	1.05	0.57	0.78	
11/1/00						0.40									
11/15/00						0.49	0.40		0.66	0.72		0.78	0.72	0.74	
11/29/00					0.24	0.13	0.19	0.11	0.17	0.08	0.28	0.10	0.09	0.08	
12/11/00					0.26		0.32	0.37	0.43	0.29		0.00		0.54	0.01
12/15/00	0.78			0.65	ŀ							0.36	1.09	0.54	0.81
12/18/00	 			0.67	l				1		0.50	0.00	0.54	0.10	
1/8/01	١		0.50	0.40		0.55	0.51	0.40	0.50	0.57	0.50	0.80	0.54	0.18	
1/22/01	1.17		0.52	0.48	0.40	0.55	0.51	0.48	0.56	0.56	0.62	0.60	0.71	0.64	
2/5/01	2.16		0.48	0.46	0.40	0.35	0.58	0.46	0.87	0.77	0.86	0.74	1.08	0.94	
2/19/01	2.30		0.80	0.42	0.73	0.61	0.55	0.46	0.70	0.71	0.64	0.62	0.95	0.73	
3/7/01	1.37		0.19	0.17	0.29	0.20	0.39	0.25	0.24	0.17	0.16	0.13	0.21	0.22	
3/19/01	2.30		0.64	0.66	0.86	0.75	0.88	0.77	0.74	0.70	0.75	0.72	0.71	0.70	

Ammonia-N (mg/L) (field)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
3/2/00			•										0.24	0.48	
3/7/00	0.24	0.84	1.20	0.48					0.60	0.72	0.60	0.48			0.96
3/23/00	0.24								0.60	0.60	0.48	0.48	0.72	0.48	0.72
3/28/00	0.48				İ				0.36	0.24	0.36	0.48	0.60	0.24	0.48
4/5/00	0.48								0.36	0.48	0.48	0.60	0.72	0.60	0.72
4/10/00	0.60				ŀ					,			0.84	0.96	
4/12/00	0.72	0.84	1.20	3.60					0.12	0.36	1.20	0.48	0.96		0.96
4/19/00									0.45	0.48	0.36	0.72	0.84	0.60	0.60
4/26/00	0.48		*						0.36	0.36	0.60	0.72	1.32	0.36	0.72
5/3/00	0.72				ļ				0.36	0.48	0.60	0.48		0.48	0.84
5/11/00	0.36	0.72	0.84	0.60					0.36	0.24	0.60	0.48	0.24	0.48	1.44
5/17/00	0.84				1				0.36	0.60	0.84	0.60	0.48	0.24	0.96
5/24/00	0.84								0.48	0.24	1.20	1.08	1.56	0.36	0.60
5/31/00	0.72								0.00	0.24	2.16	2.40	1.44	0.60	0.12
6/6/00	0.72								0.36	0.48	2.28	1.08	1.56	0.48	0.36
6/15/00	0.36	0.60	0.72	0.96					0.48	0.48	0.72	0.12	0.00	0.24	0.96
6/22/00	0.48								0.00	0.48	1.08	0.78	1.68	0.60	0.48
6/28/00	0.72								0.48	0.00	1.44	0.72	1.92	0.48	0.48
7/5/00	0.96				1				0.60	0.48	1.20	1.44	2.04	0.60	0.60
7/13/00	0.96						٠		0.00	0.00	1.20	1.68	1.56	0.72	0.72
7/21/00	1.08				l				0.72	0.72	1.92	1.80	1.68	0.60	0.84
7/24/00	0.84	0.84	0.60	0.72					0.72	0.60	0.96	0.60	0.60	0.72	1.08
8/2/00	0.84								0.60	0.60	0.96	0.72	1.68	0.60	0.84
8/9/00	0.84		•		0.48		0.96		0.60	0.72	0.84	0.60	2.04	0.72	0.84
8/16/00	1.08				0.60	0.72	0.96	0.84	0.60	0.60	0.72	0.84	2.16	0.72	0.84
8/21/00	0.48	0.60	0.72	0.72	0.48	0.60	0.96	0.84	0.12	0.48	0.24	0.12	0.84	0.36	1.08
8/31/00	0.84				0.84	0.96	0.84	1.08	0.36	0.60	0.48	0.48	0.72	0.48	0.72
9/6/00	0.48				0.72	0.72	0.84	0.96	0.60	0.36	0.36	0.24	0.84	0.72	0.96
9/13/00	0.48				0.60	0.96	0.96	0.96	0.60	0.48	0.60	0.72	0.96	0.60	0.96
9/19/00	0.48	0.48	1.44	1.20	0.84	0.96	0.96	0.96	0.60	0.60	0.72	0.48	0.72	0.48	0.96
9/26/00	0.24				0.84	0.96	1.20	1.44	0.48	0.60	0.36	0.48	0.72	0.36	1.20
10/4/00	0.48				0.72	0.84	0.62	0.84	0.36	0.48	0.72	0.48	1.20	0.72	0.96
10/18/00		0.72			0.72	0.84	0.96	0.96	0.60	0.60	0.48	0.60	0.48	0.60	
11/1/00	l				0.96	1.08	1.08	1.08	0.48	0.72	0.84	0.60	1.80	0.60	
11/15/00		0.60			0.84	0.72	0.96	0.96	0.36	0.48	0.48	0.60	0.48	0.36	
11/29/00	1				1.08	0.48	0.84	1.20	0.84	0.60	0.72		1.92	0.48	
12/11/00	1	0.60			0.84	0.96	1.44	1.60	0.60	0.48					
12/15/00	0.48				}				1		0.60	0.72	0.60	0.48	0.96
12/18/00			3.60	3.60	1				1				ĺ		
1/8/01					1								ĺ		
1/22/01	l								İ				ĺ		
2/5/01									1				l		
2/19/01	l ·														
3/7/01									1				1		
3/19/01					1				l			•	l		

CO₂ (mg/L) (field)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
3/2/00											-		10	8	
3/7/00	56	40	22						38	24	30	36			48
3/23/00	44								32	16	22	i	28	48	8
3/28/00	32				i				28	12	20	-	18	34	10
4/5/00	40				ļ				60	15	15	ļ	20	35	10
4/10/00	40								ŀ	•			30	50	
4/12/00	50	15	35						15	15	20	10	15		40
4/19/00									45	25	30		30	45	15
4/26/00.	65								35	35	45	-	25	55	20
5/3/00	30								40	30	45			45	25
5/11/00	40	20	40		İ				30	35	35	50	25	25	45
5/17/00	30								35	30	25		25	35	30
5/24/00	36			`					28	22	32	- 1	20	38	10
5/31/00	40					-			45	35	30	1	20	50	0
6/6/00	60				}				50	30	35	1	20	45	0
6/15/00	35	45	55						40	40	45	55	60	70	45
6/22/00	40								50	45	50		25	50	0
6/28/00	40				1				55	35	45		15	50	Ö
7/5/00	45								55	50	45		20	65	Ö
7/13/00	38								64	42	46		24	58	Ö
7/21/00	30								30	20	25		25	35	Ö
7/24/00	40	35	40						30	45	30	45	55	45	35
8/2/00	40							*	40	45	45	.	20	45	0
8/9/00	35			20		40			45	45	35		. 25	40	Ö
8/16/00	35			20	20	40	30		50	40	30		20	50	0
8/21/00	30	30	30	25	25	40	35		25	30	35	35	30	30	35
8/31/00	25			30	25	45	40		30	20	25	55	25	30	20
9/6/00	25			30	25	30	30		25	25	. 20	- 1	20	20	25
9/13/00	25			30	30	25	35		30	20	25	- 1	20	30	25
9/19/00	25	35	40	25	30	35	30		35	30	35	30	20	25	30
9/26/00	20			45	40	40	35		35	25	35	- 50	30	35	35
10/4/00	30			30	35	40	30		30	30	30		15	20	25
10/18/00	t	50		40	35	50	50		40	35	35		40	25	23
11/1/00				55	45	50	45		35	35	40		25	40	
11/15/00	-	10		25	30	20	35		35	20	15		20	20	
11/29/00				45	40	55	- 55		25	35	-		20	30	
12/11/00		25		35	30	30	30		35						
12/15/00										25	15	i	10	15	25
		-	60		ł				Ì		20	60	20	10	22
									l						
	1														
]						
					1										
12/18/00 1/8/01 1/22/01 2/5/01 2/19/01 3/7/01 3/19/01			60									60			

PO₄ (mg/L) (field)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
3/2/00			•										0.32	0.18	
3/7/00	0.00	0.00	0.12	0.00					0.00	0.00	0.00	0.00			0.00
3/23/00	0.00								0.10	0.10	0.00	0.12	0.00	0.88	0.10
3/28/00	0.10			;					0.00	0.08	0.00	0.00	0.00	0.00	0.00
4/5/00	0.00								0.20	0.10	0.10	0.10	0.00	0.10	0.10
4/10/00	0.00				İ	•			ŀ				0.10	0.00	-
4/12/00	0.10	1.00	0.10	1.00	İ				0.10	0.00	0.60	0.00	0.00		0.00
4/19/00									0.10	0.10	0.00	0.10	0.00	0.00	0.10
4/26/00	0.10								0.10	0.10	0.00	0.00	0.00	0.10	0.10
5/3/00	0.00								0.10	0.00	0.00	0.00		0.00	0.10
5/11/00	0.00	0.00	0.10	0.10					0.00	0.00	0.00	0.10	0.00	0.00	0.10
5/17/00	0.20								0.00	0.10	0.00	0.00	0.10	0.10	0.00
5/24/00	0.20								0.10	0.00	0.00	0.00	0.00	0.10	0.10
5/31/00	0.20								0.20	0.10	0.15	0.15	0.05	0.10	0.00
6/6/00	0.20								0.10	0.00	0.10	0.10	0.00	0.10	0.00
6/15/00	0.20	0.10	0.10	0.10					0.10	0.10	0.10	0.18	0.10	0.16	0.12
6/22/00	0.30								0.10	0.30	0.20	0.20	0.00	0.10	0.00
6/28/00	0.20								0.10	0.05	0.10	0.05	0.05	0.10	0.00
7/5/00	0.30								0.05	0.10	0.10	0.10	0.00	0.10	0.00
7/13/00	0.30				ĺ				0.10	0.20	0.30	0.20	0.00	0.10	0.00
7/21/00	0.20								0.10	0.10	0.20	0.20	0.10	0.10	0.10
7/24/00	0.18	0.00	0.08	0.12					0.10	0.08	0.10	0.10	0.10	0.14	0.12
8/2/00	0.20								0.10	0.10	0.00	0.10	0.10	0.00	0.10
8/9/00	0.10			3	0.00		0.04		0.12	0.10	0.04	0.08	0.00	0.00	0.00
8/16/00															
8/21/00	0.10	0.10	0.20	0.10	0.10	0.10	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.10	0.10
8/31/00	0.20				0.00	0.10	0.10	0.00	0.10	0.10	0.10	0.10	0.00	0.10	0.00
9/6/00	0.05				0.05	0.10	0.05	0.05	0.15	0.10	0.05	0.10	0.10	0.05	0.05
9/13/00	0.10				0.10	0.05	0.05	0.05	0.10	0.10	0.05	0.10	0.10	0.05	0.05
9/19/00	0.05	0.10	0.20	0.40	0.10	0.10	0.10	0.01	0.10	0.15	0.05	0.05	0.50	0.10	0.05
9/26/00	0.00				0.10	0.00	0.00	0.00	0.10	0.00	0.10	0.00	0.10	0.00	0.00
10/4/00	0.10				0.10	0.00	0.00	0.00	0.10	0.00	0.10	0.10	0.10	0.00	0.00
10/18/00		0.00			0.00	0.00	0.10	0.05	0.05	0.05	0.05	0.05	0.10	0.05	
11/1/00					0.10	1.60	0.00	0.00	0.50	0.00	0.10	0.10	0.00	0.00	
11/15/00		0.00			0.10	0.00	0.10	0.00	0.00	0.00	0.10	0.00	0.00	0.10	
11/29/00	ŀ				0.00	0.00	0.00	0.00	0.40	0.00	0.00		0.10	0.00	
12/11/00		0.05			0.00	0.00	0.00	0.00	0.08	0.02					
12/15/00	0.05										0.10	0.05	0.00	0.10	0.05
12/18/00	ŀ		15.00	16.60					}						
1/8/01									1						
1/22/01	l														
2/5/01	l														
2/19/01	l														
3/7/01	1														
3/19/01	Ī				ļ.				Į.						

Laboratory Data

Ammonia-N (mg/L) (laboratory)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
Mar-00	<0.01	0.19	0.15	0.04					0.11	0.10	0.06	0.07	0.02	0.03	0.3
Apr-00	<0.05	0.21	0.39	3.10					0.02	0.02	0.06	0.05	<0.01	< 0.01	0.27
May-00	< 0.01	0.19	0.17	0.18					<0.01	< 0.01	0.16	<0.01	<0.01	< 0.01	0.26
Jun-00	< 0.01	0.19	0.17	0.21					<0.01	0.04	0.18	0.02	<0.01	< 0.01	0.31
Jul-00	<0.01	0.20	0.12	0.11					<0.01	< 0.01	0.37	0.10	<0.01	0.01	0.26
Aug-00	< 0.01	0.10	0.11	0.10	0.04	0.04	0.15	0.15	<0.01	< 0.01	0.10	<0.01	0.02	0.10	0.21
Sep-00	< 0.01	0.17	0.44	0.04	0.17	0.17	0.23	0.24	<0.01	< 0.01	< 0.01	<0.01	<0.01	< 0.01	0.28
Oct-00		0.08			0.14	0.13	0.17	0.18	0.05	< 0.01	< 0.01	<0.01	<0.01	< 0.01	
Nov-00		0.07			0.14	0.14	0.22	0.23	<0.01	< 0.01	<0.01	<0.01	<0.01	< 0.01	
Dec-00	0.02	0.07	14	15	0.18	0.18	0.44	0.41	<0.01	< 0.01	< 0.01	< 0.01	<0.01	0.03	0.26

PO₄ (mg/L) (laboratory)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
Mar-00	0.02	0.02	0.02	0.04					0.01	0.01	0.02	0.01	0.08	0.19	0.05
Apr-00	0.04	0.01	0.13	0.90					0.04	0.04	0.02	0.03	0.06	0.27	0.05
May-00	0.01	0.03	0.02	0.02					0.01	0.02	0.005 U	0.01	0.02	0.03	0.01
Jun-00	< 0.021	< 0.017	0.03	0.04	ł				<0.011	< 0.012	<0.019	< 0.03	<0.012	< 0.022	0.04
Jul-00	0.019 J	0.02 J	0.043 J	0.032 J	İ				0.01 J	0.01 J	0.087 J	0.04 J	0.023 J	0.019 J	0.02 J
Aug-00	0.03	0.05	0.09	0.04	0.05	0.05	0.02	0.02	0.03	0.02	0.03	0.03	0.02	0.02	0.03
Sep-00	0.03	0.03	0.14	0.73	0.02	0.01	0.01	0.01	0.02	0.03	0.01	0.04	0.02	0.04	0.02
Oct-00		0.01			0.02	0.02	0.03	0.04	0.03	0.02	0.06	0.02	0.02	0.03	
Nov-00		0.01			0.01	0.01	0.01	0.01	0.03	0.02	0.02	0.02	0.02	0.03	
Dec-00	0.05	0.01	14.00	15.00	0.01	0.01	0.01	0.01	0.02	0.02	0.02	0.02	0.01	0.02	0.03

Chloride (mg/L) (laboratory)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
Мат-00	4.9	9.1	7.9	5.4			٠.		4.9	4.7	5.5	4.9	5.0	14.0	5.6
Apr-00	4.8	7.4	6.8	6.9	l				5.8	5.7	4.6	4.4	5.0	13.0	5.4
May-00	4.8	7.7	7.0	6.9					6.4	6.6	5.8	6.2	6.2	11.0	5.8
Jun-00	4.5	7.1	6.5	6.2					5.8	6.3	6.1	6.2	6.0	8.1	5.6
Jul-00	4.7	7.5	6.5	6.7					6.1	6.3	6.5	6.4	5.6	11.0	5.6
Aug-00	5.3	7.6	7.0	6.7	7.2	7.2	7.1	7.1	6.2	6.5	6.8	6.4	6.3	10.0	5.2
Sep-00	5 J	8 J	7.4 J	7.6 J	7.6 J	7.6 J	7.2 J	7.5 J	6.8 J	7.2 J	6.6 J	6.8 J	6.6 J	12 J	5.7 J
Oct-00		7.1			7.5	7.5	7.1	7.2	6.8	7.3	6.8	6.7	6.7	9.6	
Nov-00		6.8 J			7.2 J	7.1 J	7 J	7.1 J	6.7 J	7 J	6.7 J	6.6 J	6.4 J	9.9 J	
Dec-00	4.8	6.9	9.5	9.6	7.8	7.4	8.3	8.1	6.6	7.1	7.0	7.1	6.8	11.0	5.4

TOC (mg/L) (laboratory)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
Mar-00	<1	4.7	1.7	6.2					<1	<1	4.2	2.2	<1	<1	<1
Apr-00	<1	<1	<1	56	İ				<1	<1	• 1	<1	<1	<1	<1
May-00	<1	1.8	<1	<1					<1	<1	1.6	<1	<1	<1	<1
Jun-00	<1	<1	<1	<1					<1	<1	<1	<1	<1	<1	<1
Jul-00	<1	<1	<1	<1					<1	<1	<1	<1	<1	<1	<1
Aug-00	<1	<1	<1	3.1	6.5	13	16	22	1.3	<1	1.1	<1	<1	<1	1.3
Sep-00	<1	<1	5.7	100	14	8.9	7	8.5	<1	<1	<1	<1	<1	<1	<1
Oct-00		1.1			7	10	5.5	6.9	<1	<1	<1	1.1	<1	1.5	
Nov-00		<1			9.2	9.1	77	78	1	1.2	1.6	1	1.7	2.8	
Dec-00	1	1.1	1,000	1,300	140	110	380	390	1	1.4	<1	<1	1.3	1.8	<1

NO₃ (mg/L) (laboratory)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
Mar-00	1.1	< 0.2	< 0.2	0.3					0.8	0.8	< 0.2	< 0.2	1.6	3.2	< 0.2
Apr-00	1.6	0.2	< 0.2	< 0.2	1				1.4	1.4	< 0.2	0.4	1.2	3.4	< 0.2
May-00	1.0	0.3	0.6	0.6					0.8	0.7	0.6	0.8	1.2	2.2	< 0.2
Jun-00	1.0	0.8	0.6	0.5					0.6	0.5	0.5	0.6	0.8	1.2	< 0.2
Jul-00	0.9	0.7	0.7	0.7					< 0.2	0.5	0.5	0.4	0.5	2.5	< 0.2
Aug-00	0.9	0.6	< 0.2	0.62	< 0.2	< 0.2	< 0.2	< 0.2	0.41	0.34	0.48	0.51	0.5	1.9	< 0.2
Sep-00	0.9	0.3	< 0.2	< 0.2	< 0.2	< 0.2	< 0.2	< 0.2	0.21	< 0.2	0.29	< 0.2	0.3	2.1	. < 0.2
Oct-00		0.2			< 0.2	< 0.2	< 0.2	< 0.2	0.22	< 0.2	0.31	0.27	0.3	1.4	
Nov-00		0.6			<0.2	< 0.2	< 0.2	< 0.2	< 0.2	< 0.2	0.23	0.2	0.2	2.0	
Dec-00	0.9	0.5	< 0.2	< 0.2	<0.2	< 0.2	< 0.2	<0.2	<0.2	< 0.2	< 0.2	< 0.2	0.2	2.9	< 0.2

TKN (mg/L) (laboratory)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
Mar-00	0.6 J	0.9 J	0.6 J	0.8 J				•	0.7 J	0.8 J	0.5 J	<0.3	0.7 J	0.8 J	1 J
Apr-00	< 0.3	0.3	1.2	6.5					0.5	< 0.3	0.4	0.4	< 0.3	0.3	0.6
May-00	0.3	0.6	0.6	0.6					0.4	0.6	0.8	< 0.3	< 0.3	0.7	0.6
Jun-00	0.6	1.1	1.9	7.5					0.4	0.3	0.9	0.4	0.6	0.9	1.3
Jul-00	1	0.8	1	1.7	l				0.4	0.8	1.3	0.5	< 0.5	1.3	1.1
Aug-00	0.34	2.2	0.61	0.56	0.6	0.7	0.7	1	< 0.3	< 0.3	0.47	< 0.3	0.37	1.6	0.78
Sep-00	0.3	1.1	1.5	2.4	0.9	0.9	0.9	0.9	0.5	0.4	0.7	0.6	0.5	0.3	1
Oct-00		< 0.4			< 0.4	< 0.5	< 0.6	< 0.8	< 0.3	< 0.4	< 0.06	< 0.6	< 0.3	< 0.5	
Nov-00		< 0.3			< 0.3	< 0.3	0.5	0.5	<0.3	< 0.3	< 0.3	< 0.3	< 0.3	< 0.3	
Dec-00	0.5	< 0.3	14	16	< 0.3	0.3	0.8	0.8	<0.3	< 0.3	< 0.3	< 0.3	< 0.3	< 0.3	0.5

Total P (mg/L) (laboratory)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
Mar-00	0.05 J	0.08 J	0.07 J	0.09 J					0.04 J	0.04 J	0.05 J	0.06 J	0.07 J	0.06 J	0.11 J
Apr-00	0.07 J	0.1 J	0.37 J	5.4 J	ĺ				0.08 J	0.05 J	0.03 J	0.06 J	0.13 J	0.11 J	0.04 J
May-00	0.1 J	0.18 J	0.16 J	0.04 J					0.03 J	0.04 J	0.03 J	0.06 J	0.1 J	0.06 J	0.02 J
Jun-00	0.04	0.12	0.08	0.11	1				0.06	0.04	0.11	0.03	0.15	0.09	0.09
Jul-00	0.19 J	0.12 J	0.25 J	0.25 J	ŀ				0.16 J	0.15 J	0.1 J	0.19 J	0.14 J	0.1 J	0.05 J
Aug-00	0.099	0.4	0.095	0.13	0.053	0.054	0.041	0.18	0.21	0.053	0.063	0.086	0.057	0.09	0.059
Sep-00	0.06 J	0.27 J	0.2 J	0.86 J	0.031 J	0.028 J	0.032 J	0.075 J	0.074 J	0.1 J	0.047 J	0.23 J	0.049 J	0.06 J	0.036 J
Oct-00		0.025			0.034	0.046	0.13	0.077	0.039	0.13	0.18	0.21	0.029	0.11	
Nov-00		0.21 J			<0.02	< 0.02	< 0.02	< 0.02	< 0.02	< 0.02	< 0.02	< 0.02	<0.02	< 0.02	
Dec-00	0.04	< 0.02	16	<2	0.03	0.02	0.08	0.11	0.1	4	0.3	0.1	0.03	0.02	0.08

HPC (CFU/mL) (laboratory)

Date	3LF	57LF	60LF	61LF	62LFA	62LFB	63LFA	63LFB	58LFA	58LFB	59LFA	59LFB	4LF	4UF	21LF
Mar-00	9.8E+02	2.4E+05	1.6E+05	5.4E+04	,				1.5E+04	2.1E+04	1.5E+05	1.2E+05			1.9E+04
Apr-00	6.0E + 03	1.4E+05	2.5E+06	1.7E+06					6.1E+03	5.9E+03	3.2E+05	1.7E+05	1.3E+04	1.5E+04	3.5E + 02
May-00	7.4E + 03	1.4E+06	1.0E+05	6.1E+04					1.4E+04	8.5E + 02	1.4E+05	2.0E + 04	1.4E+03	9.2E + 02	2.8E+02
Jun-00	1.3E+03	1.7E+05	8.4E+04	4.0E+05				•	1.5E+03	5.2E+02	2.1E+04	9.2E+03	7.4E+03	6.4E+03	4.3E+02
Jul-00	4.3E+03	3.1E+04	1.5E+04	8.1E+04					1.1E+04	6.0E+02	5.2E+03	2.9E+03	2.5E+04	3.4E+03	2.5E + 02
Aug-00	1.6E+03	1.2E+05	1.2E+05	9.6E+04	8.8E+03	5.6E+03	2.3E+04	1.8E+04	2.9E+03	7.0E+02	3.7E+04	1.1E+03	5.4E+03	7.8E+03	2.6E+04
Sep-00	1.1E+04	5.8E+04	7.9E + 04	8.1E+06	3.1E+03	6.8E+03	4.1E+03	1.3E+03	2.1E+04	3.1E+02	1.9E+04	3.2E+03	4.1E+04	3.1E+04	1.9E+04
Oct-00		2.6E+03			6.9E+02	6.0E + 02	2.0E+03	1.0E+02	1.5E+03	5.9E+03	8.1E+04	2.8E+02	4.4E+03	2.7E + 03	
Nov-00		2.2E + 03			5.7E+02	1.5E+02	1.6E+03	5.2E+02	1.8E+03	1.0E+02	2.4E + 04	1.4E+04	3.6E+03	1.1E+03	
Dec-00	3.6E+02	1.2E+03	2.8E+05	8.2E+05	1.7E+03	1.3E+03	1.1E+03	4.0E+02	5.5E+03	1.9E+02	3.2E+04	1.6E+03	8.5E+03	1.8E+03	5.3E+03
Mean	4.1E+03	2.2E+05	4.2E+05	1.4E+06	3.0E+03	2.9E+03	6.4E+03	4.1E+03	8.0E+03	3.6E+03	8.3E+04	3.4E+04	1.2E+04	7.8E+03	8.8E+03

APPENDIX E GROUNDWATER ELEVATIONS

Groundwater Elevations							
Well	TOC Elev (msl)	Date	DTW (ft)	Elev (msl)			
007G03LF	283.32	3/7/00	29.40	253.92			
		3/14/00	29.44	253.88			
		3/23/00	29.09	254.23			
		4/10/00	28.87	254.45			
	,	5/11/00	28.40	254.92			
		6/15/00	28.40	254.92			
		7/16/00	28.90	254.42			
		8/21/00	29.85	253.47			
	1	9/19/00	. 30.44	252.88			
		10/18/00	30.94	252.38			
l		11/15/00	31.17	252.15			
		12/12/00	31.18	252.14			
007G04LF	283.12	3/7/00	29.47	253.65			
		3/14/00	29.58	253.54			
		3/23/00	29.53	253.59			
		4/10/00	28.82	254.3			
		5/11/00	28.71	254.41			
	1	6/15/00	28.91	254.21			
		7/16/00 ·	29.09	254.03			
		8/21/00	30.14	252.98			
		9/19/00	30.82	252.3			
		10/18/00	31.17	251.95			
		11/15/00	31.45	251.67			
		12/12/00	31.51	251.61			
007G04UF	283.21	3/7/00	29.51	253.7			
		3/14/00	29.71	253.5			
	1	3/23/00	29.54	253.67			
	İ	4/10/00	28.91	254.3			
	<u> </u>	5/11/00	28.72	254.49			
		6/15/00	28.66	254.55			
		7/16/00	29.17	254.04			
		8/21/00	30.15	253.06			
	1	9/19/00	30.83	252.38			
		10/18/00	31.26	251.95			
		11/15/00	31.49	251.72			
		12/12/00	31.59	251.62			
007G21LF	283.66	3/7/00	32.6	251.06			
	· .	3/14/00	32.82	250.84			
		3/23/00	32.75	250.91			
		4/10/00	32.41	251.25			
	·	5/11/00	32.22	251.44			
ŀ		6/15/00	32.31	251.35			
		7/16/00	32.82	250.84			
		8/21/00	33.55	. 250.11			
		9/19/00	34.1	249.56			
		10/18/00	'				
		11/15/00	0.4.50				
L		12/12/00	34.73	248.93			

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Groundwater Elevations							
Well	TOC Elev (msl)	Date	DTW (ft)	Elev (msl)			
007G57LF	281.71	3/7/00 3/14/00 3/23/00	28.21 28.27	253.5 253.44			
ı		4/10/00 5/11/00	27.65 27.3	254.06 254.41			
	1	6/15/00	27.54	254.17			
		7/16/00 8/21/00	27.81 28.76	253.9 252.95			
		9/19/00 10/18/00	29.44	252.27			
		11/15/00 12/12/00					
007G58LF	283.21	3/7/00	29.61 29.61	253.6			
		3/14/00 3/23/00	29.4	253.6 253.81			
		4/10/00	29.02	254.19			
		5/11/00 6/15/00	28.55 28.87	254.66 254.34			
		7/16/00	29.2	254.01			
ļ		8/21/00 9/19/00	30.1 30.82	253.11 252.39			
		10/18/00	31.21	252.0			
		11/15/00 12/12/00	31.79 31.63	251.42 251.58			
007G59LF	283.19	3/7/00	29.57	253.62			
		3/14/00	29.47	253.72			
	1	3/23/00 4/10/00	29.22 28.86	253.97 254.33			
		5/11/00	28.6	254.53 254.59			
		6/15/00	28.72	254.47			
	· ·	7/16/00 8/21/00	29.35 30.01	253.84 253.18			
		9/19/00	30.68	252.51			
		10/18/00 11/15/00	31.14 31.48	252.05 251.71			
		12/12/00	31.44	251.75			
007G60LF	282.42	3/7/00 3/14/00	28.86 28.53	253.56 253.89			
		3/23/00					
		4/10/00 5/11/00	28.2 27.92	254.22 254.5			
		6/15/00	27.72	254.7			
		7/16/00 8/21/00	28.4 27.05	254.02 255.37			
	ļ	9/19/00	30	252.42			
	}	10/18/00 11/15/00		'			
		12/12/00	30.5	251.92			
007G61LF	282.55	3/7/00	28.89	253.66			
		3/14/00 3/23/00	28.62	253.93			
		4/10/00	28.4	254.15			
1		5/11/00 6/15/00	27.97 27.95	254.58 254.60			
		7/16/00	28.46	254.09			
		8/21/00 9/19/00	29.45	253.10 253.42			
		10/18/00	30.12	252.43			
		11/15/00 12/12/00	30.5	252.05			
IL_,		12/12/00	30.3	232.03			

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Groundwater Elevations							
Well	TOC Elev (msl)	Date	DTW (ft)	Elev (msl)			
007G62LF	283.37	8/21/00 9/19/00 10/18/00 11/15/00 12/12/00	30.05 30.58 31.08 31.44 31.46				
007G63LF	283.40	8/21/00 9/19/00 10/18/00 11/15/00 12/12/00	29.88 30.4 30.92 31.38 31.3				

Notes:

TOC DTW msl

Top of (well) casing
Depth to water
Mean sea level
Depth to water measurement not taken synoptically with other wells

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APPENDIX F HYDRAULIC EVALUATION

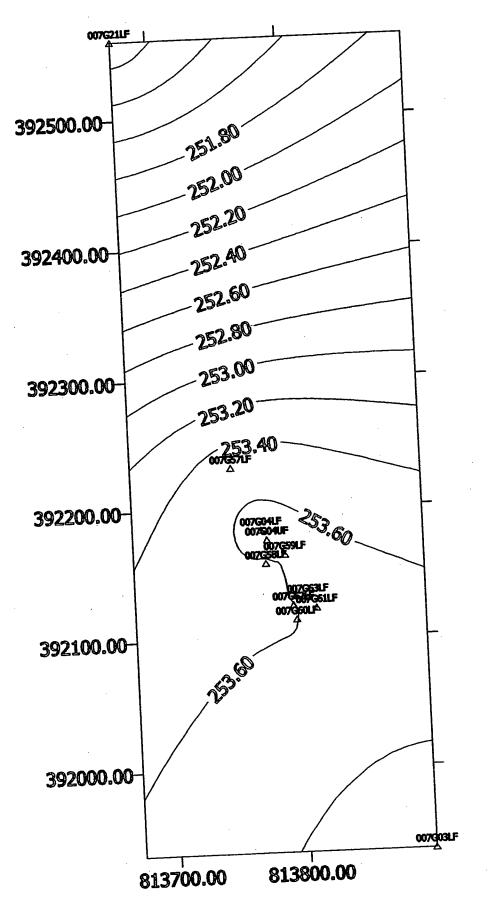
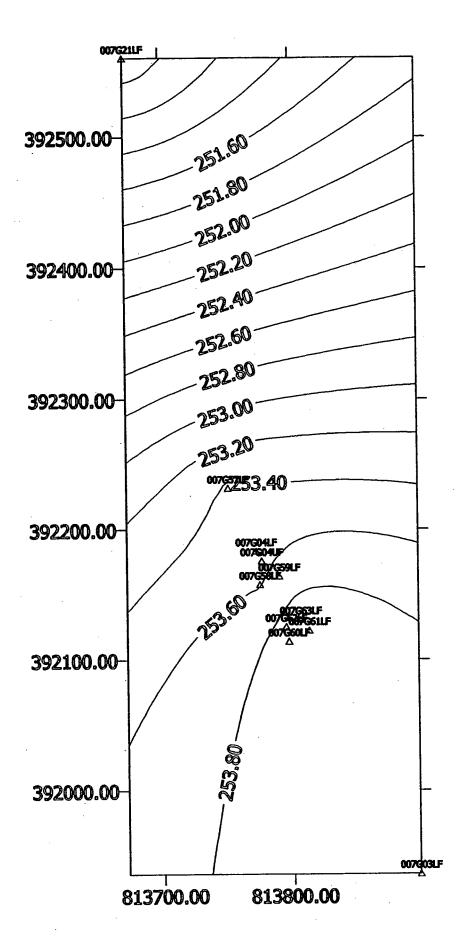


FIGURE 1
POTENTIOMETRIC SURFACE 3-8-00 (baseline)



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FIGURE 2
POTENTIOMETRIC SURFACE 3-14-00

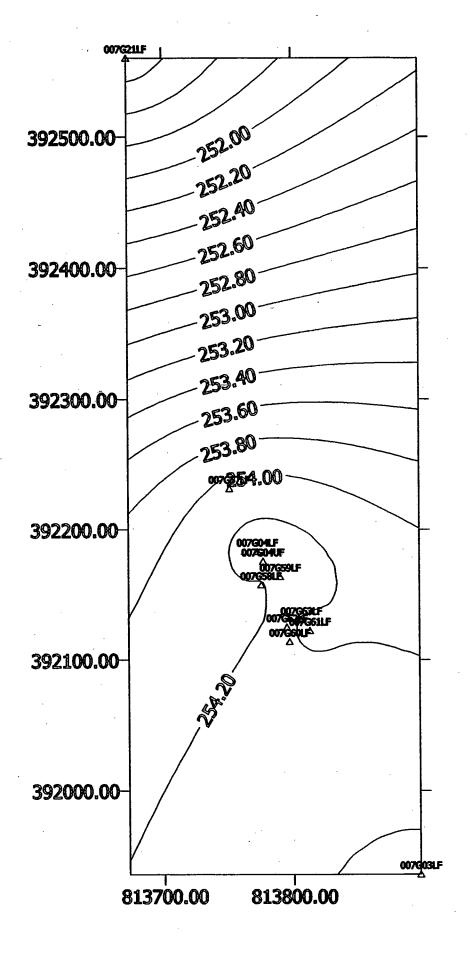


FIGURE 3
POTENTIOMETRIC SURFACE 4-10-00

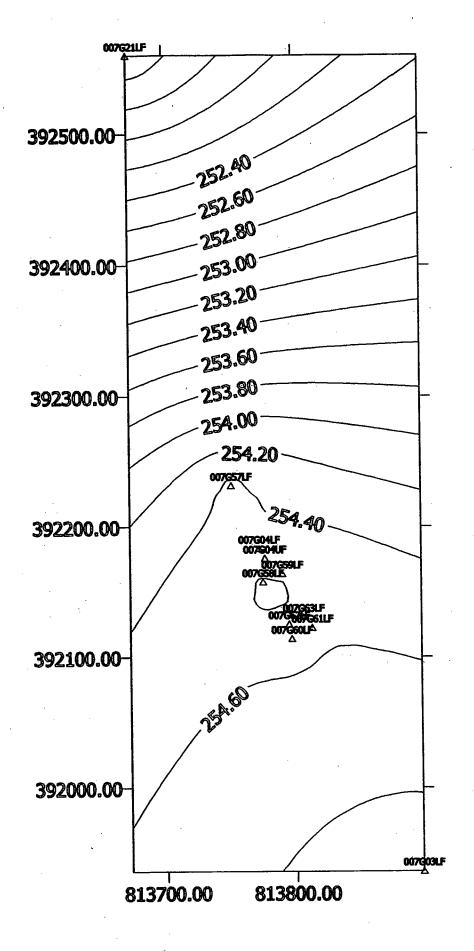


FIGURE 4
POTENTIOMETRIC SURFACE 5-11-00

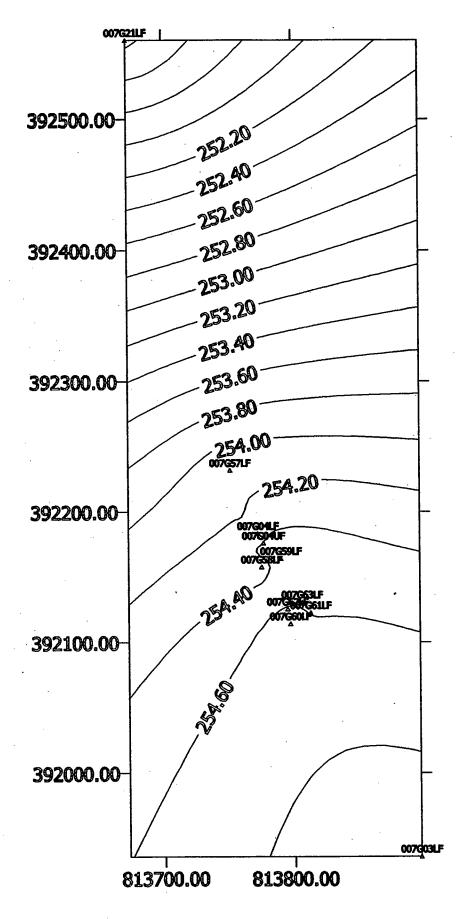


FIGURE 5
POTENTIOMETRIC SURFACE 6-15-00

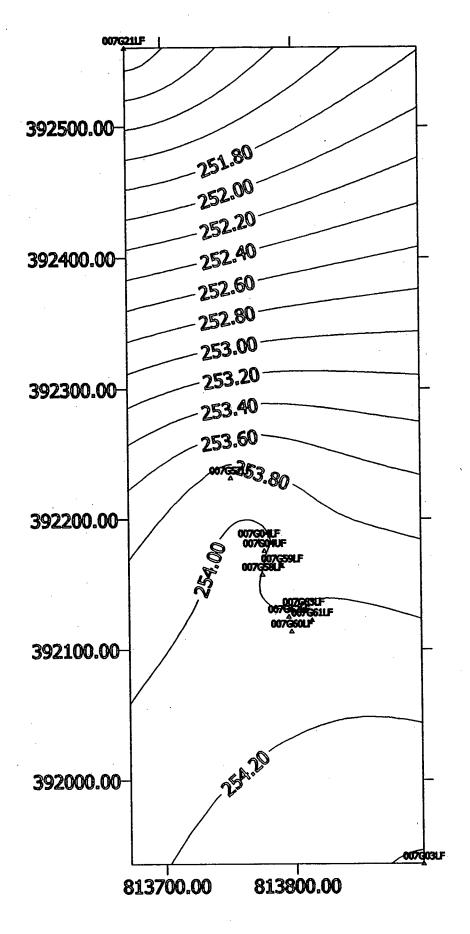


FIGURE 6
POTENTIOMETRIC SURFACE 7-16-00

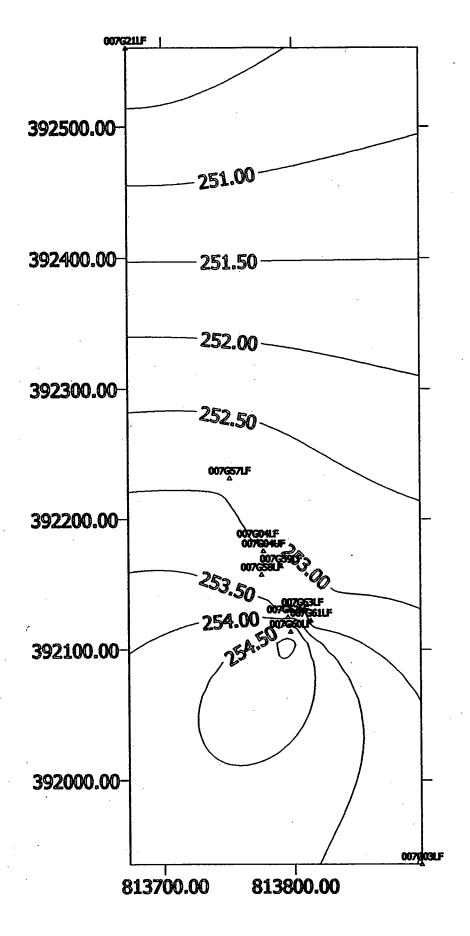


FIGURE 7
POTENTIOMETRIC SURFACE 8-21-00

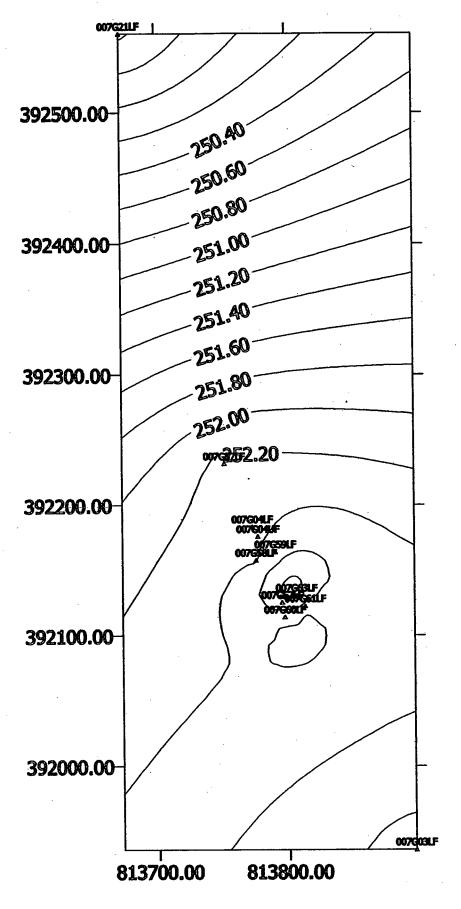


FIGURE 8
POTENTIOMETRIC SURFACE 9-19-00

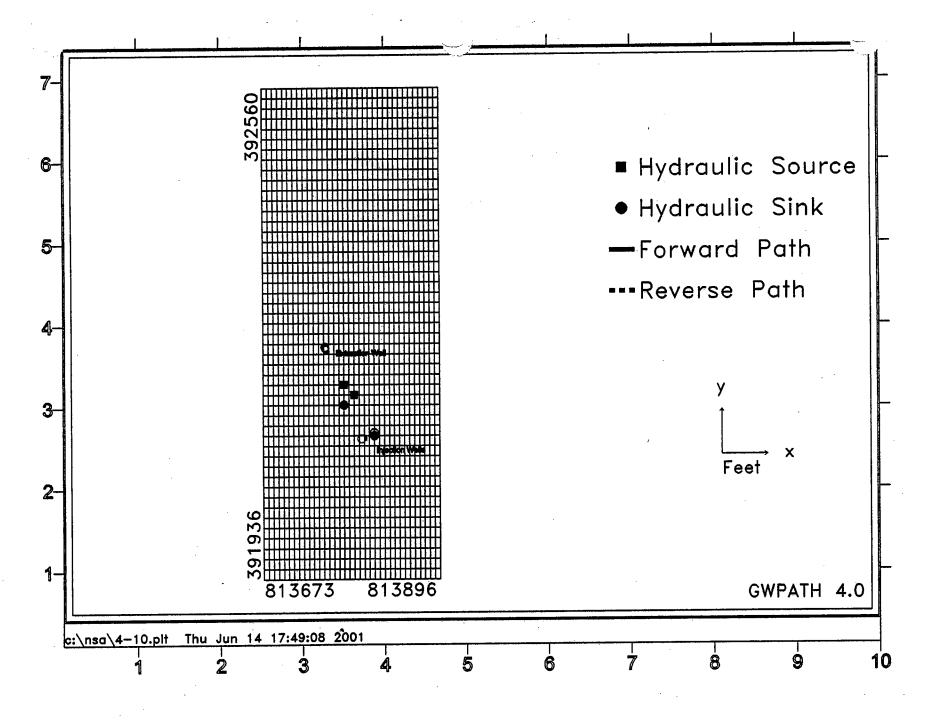


FIGURE 9 4-10-00 (~30 DAYS TRAVEL TIME)

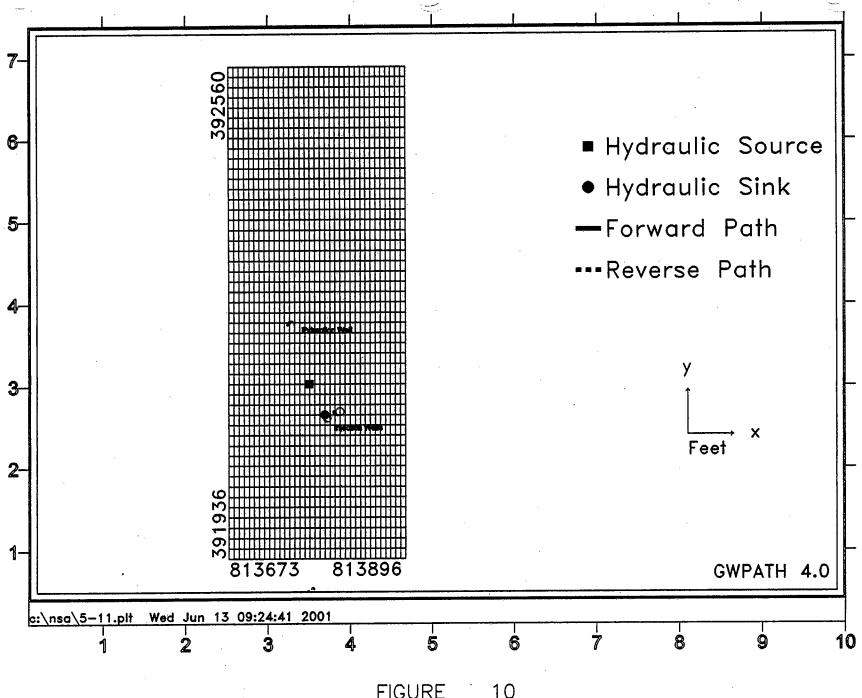


FIGURE 10 5-11-00 (~60 DAYS TRAVEL TIME)

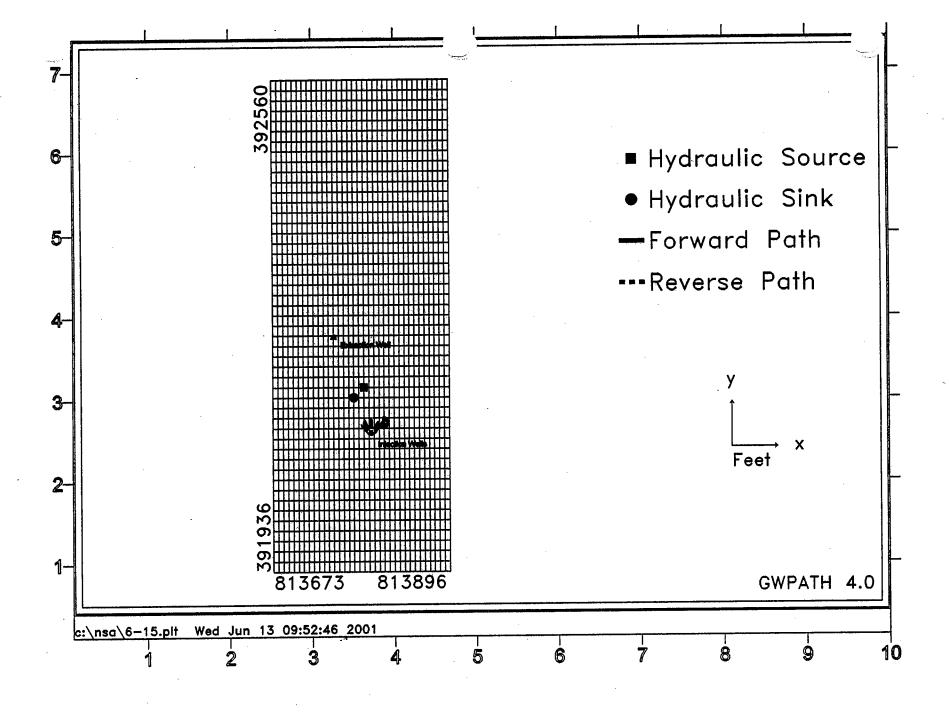


FIGURE 11 6-15-00 (~90 DAYS TRAVEL TIME)

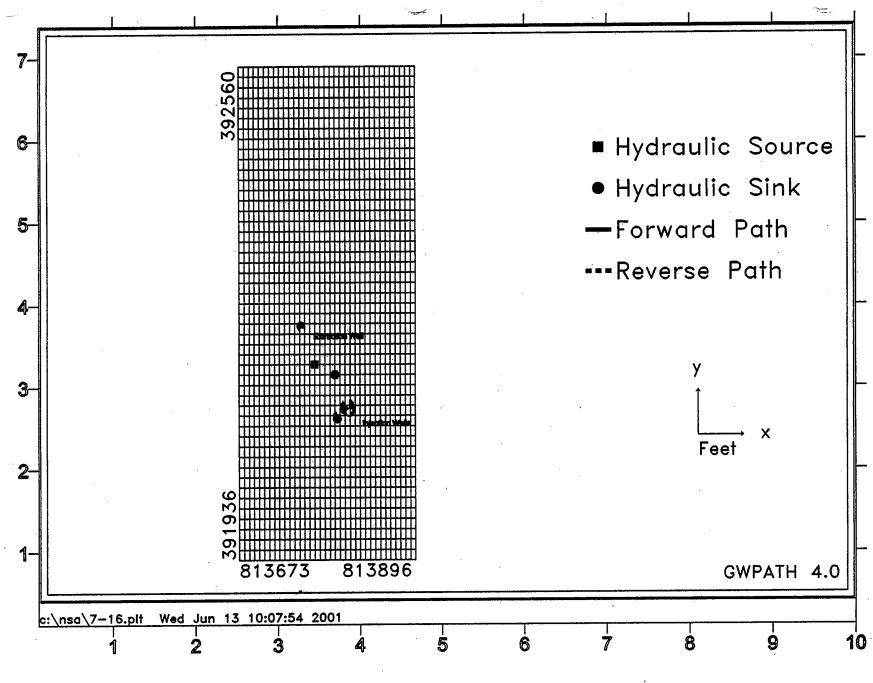


FIGURE 12 7-16-00 (~120 DAYS TRAVEL TIME)

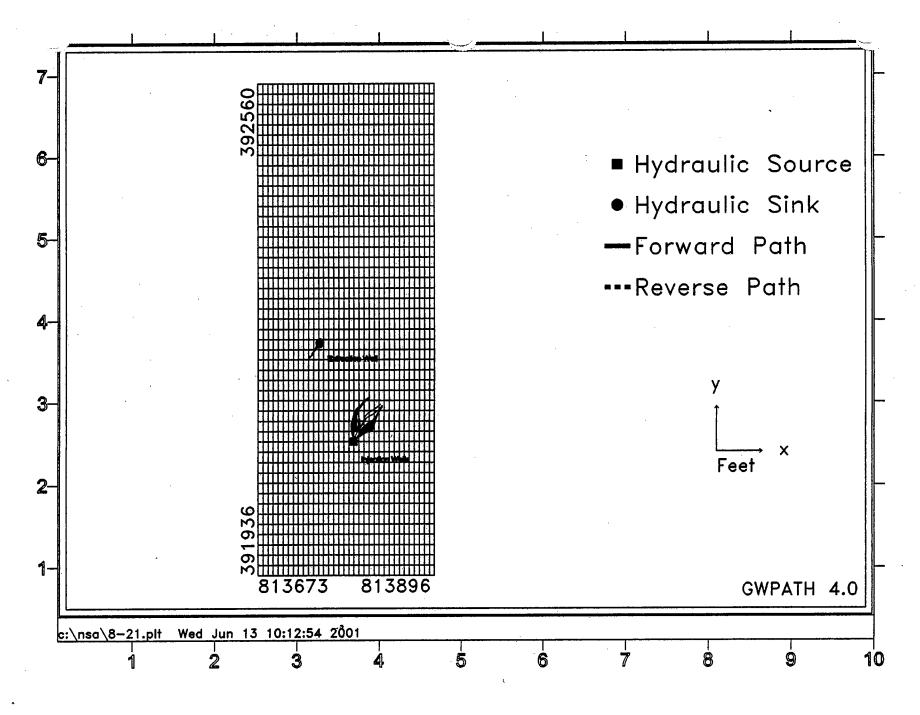


FIGURE 13 8-21-00 (~160 DAYS TRAVEL TIME)

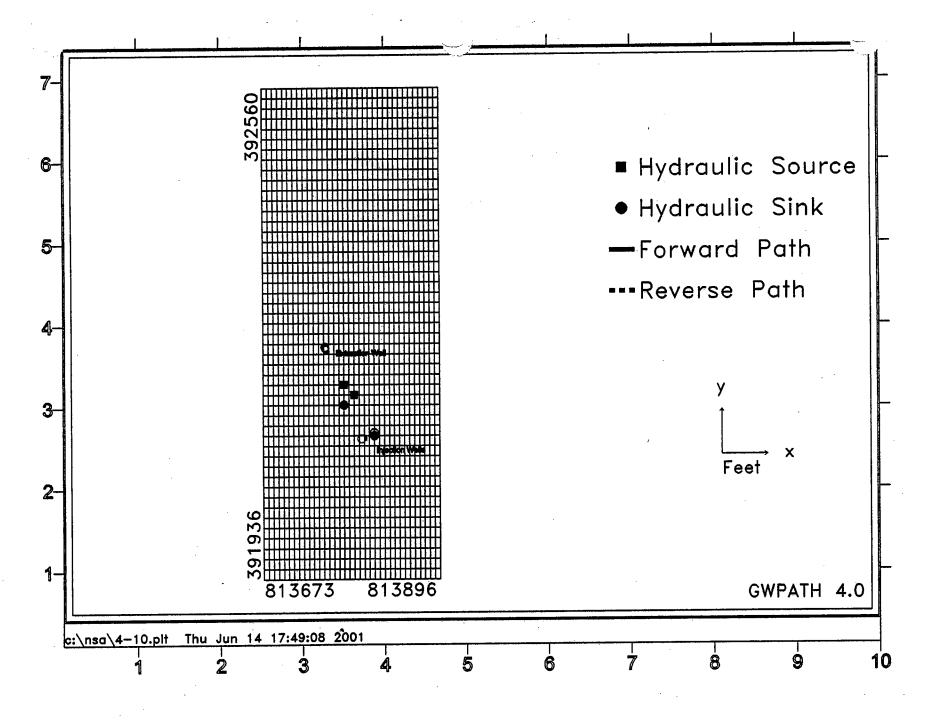


FIGURE 9 4-10-00 (~30 DAYS TRAVEL TIME)

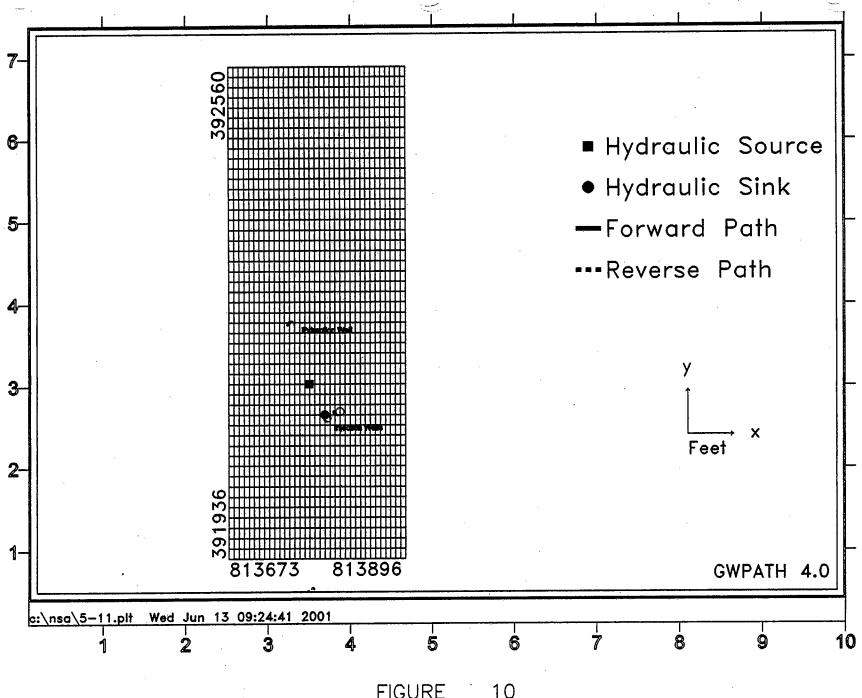


FIGURE 10 5-11-00 (~60 DAYS TRAVEL TIME)

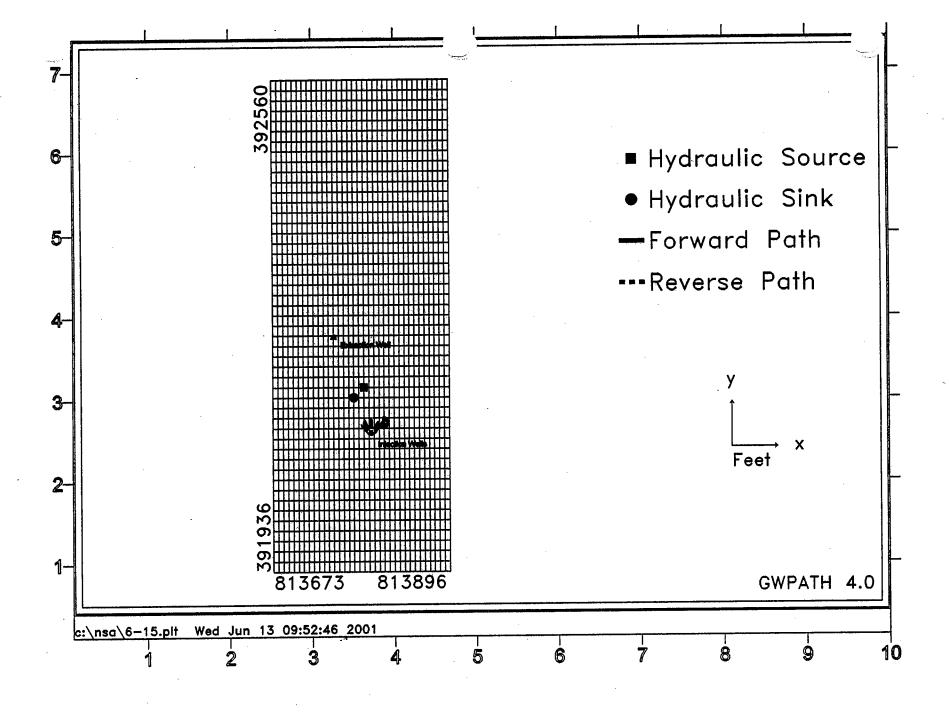


FIGURE 11 6-15-00 (~90 DAYS TRAVEL TIME)

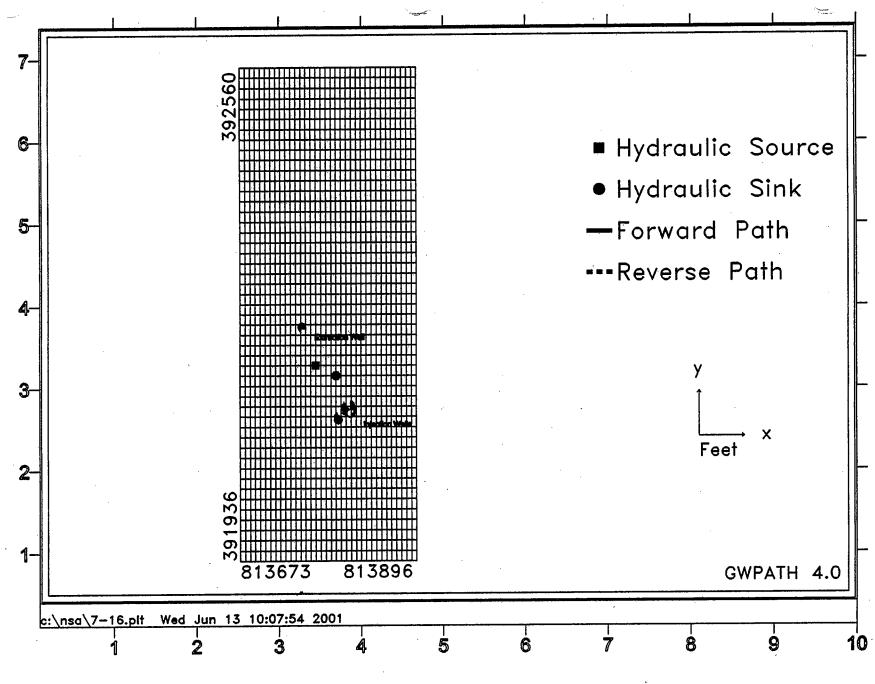


FIGURE 12 7-16-00 (~120 DAYS TRAVEL TIME)

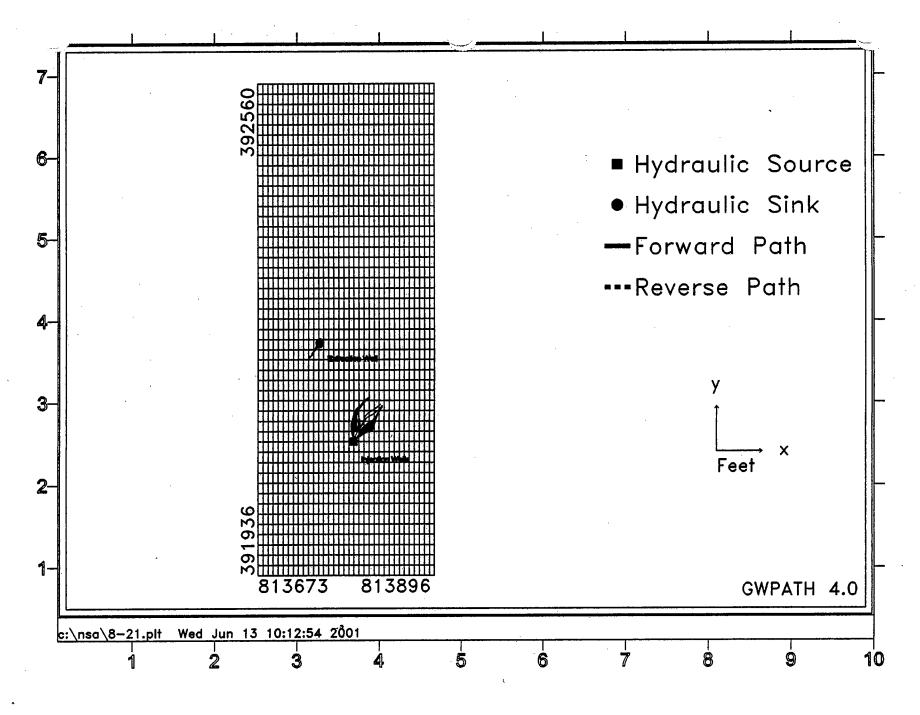


FIGURE 13 8-21-00 (~160 DAYS TRAVEL TIME)

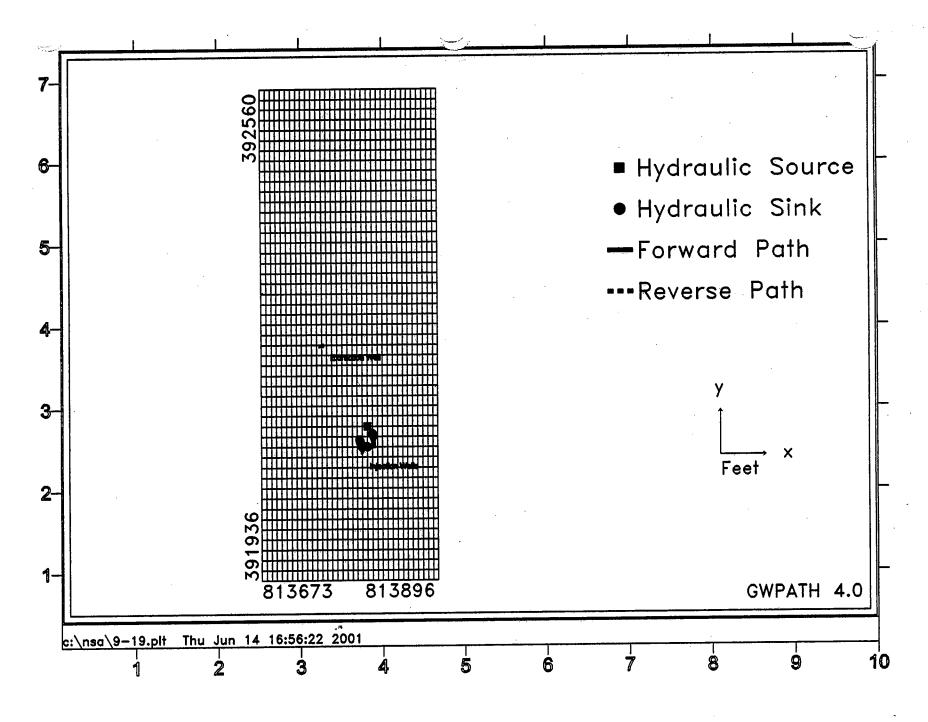


FIGURE 14 9-19-00 (~180 DAYS TRAVEL TIME)

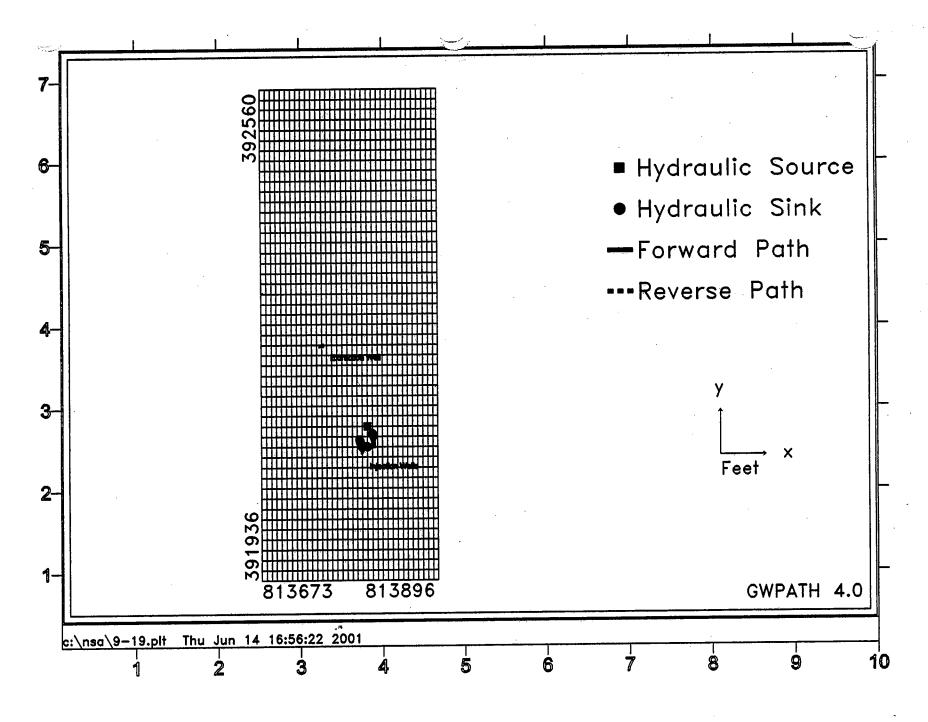


FIGURE 14 9-19-00 (~180 DAYS TRAVEL TIME)